

=> fil reg; d ide 12; d ide 13
FILE 'REGISTRY' ENTERED AT 13:26:55 ON 31 JUL 2008
USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.
PLEASE SEE "HELP USAGETERMS" FOR DETAILS.
COPYRIGHT (C) 2008 American Chemical Society (ACS)

Property values tagged with IC are from the ZIC/VINITI data file
provided by InfoChem.

STRUCTURE FILE UPDATES: 30 JUL 2008 HIGHEST RN 1037244-07-7
DICTIONARY FILE UPDATES: 30 JUL 2008 HIGHEST RN 1037244-07-7

New CAS Information Use Policies, enter HELP USAGETERMS for details.

TSCA INFORMATION NOW CURRENT THROUGH January 9, 2008.

Please note that search-term pricing does apply when
conducting SmartSELECT searches.

REGISTRY includes numerically searchable data for experimental and
predicted properties as well as tags indicating availability of
experimental property data in the original document. For information
on property searching in REGISTRY, refer to:

<http://www.cas.org/support/stngen/stndoc/properties.html>

L2 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2008 ACS on STN
RN 13981-52-7 REGISTRY
ED Entered STN: 16 Nov 1984
CN Polonium, isotope of mass 210 (CA INDEX NAME)
OTHER NAMES:
CN 210Po
CN Po 210
CN Polonium-210
CN Radium F
DR 14809-83-7
MF Po
CI COM
LC STN Files: AGRICOLA, ANABSTR, BIOSIS, BIOTECHNO, CA, CAOLD, CAPLUS,
CHEMLIST, CIN, CSNB, EMBASE, HSDB*, IFICDB, IFIPAT, IFIUDB, MEDLINE,
MSDS-OHS, PIRA, PROMT, TOXCENTER, USPAT2, USPATFULL, USPATOLD
(*File contains numerically searchable property data)

210Po

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

3023 REFERENCES IN FILE CA (1907 TO DATE)
3 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
3024 REFERENCES IN FILE CAPLUS (1907 TO DATE)
42 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

L3 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2008 ACS on STN
 RN 14255-04--0 REGISTRY
 ED Entered STN: 16 Nov 1984
 CN Lead, isotope of mass 210 (CA INDEX NAME)
 OTHER NAMES:
 CN 210Pb
 CN Lead-210
 CN Pb 210
 CN Radium D
 MF Pb
 LC STN Files: AGRICOLA, ANABSTR, BIOSIS, BIOTECHNO, CA, CAOLD, CAPLUS,
 CHEMLIST, CIN, CSNB, EMBASE, IFICDB, IFIPAT, IFIUDB, PROMT, TOXCENTER,
 USPAT2, USPATFULL, USPATOLD

210Pb

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

4375 REFERENCES IN FILE CA (1907 TO DATE)
 10 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
 4376 REFERENCES IN FILE CAPLUS (1907 TO DATE)
 21 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

=> => fil capl; d que 19; d que 116; d que 125; d que 126; d que 132; d que 135
 FILE 'CAPLUS' ENTERED AT 14:09:11 ON 31 JUL 2008
 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.
 PLEASE SEE "HELP USAGETERMS" FOR DETAILS.
 COPYRIGHT (C) 2008 AMERICAN CHEMICAL SOCIETY (ACS)

Copyright of the articles to which records in this database refer is held by the publishers listed in the PUBLISHER (PB) field (available for records published or updated in Chemical Abstracts after December 26, 1996), unless otherwise indicated in the original publications. The CA Lexicon is the copyrighted intellectual property of the American Chemical Society and is provided to assist you in searching databases on STN. Any dissemination, distribution, copying, or storing of this information, without the prior written consent of CAS, is strictly prohibited.

FILE COVERS 1907 - 31 Jul 2008 VOL 149 ISS 5
 FILE LAST UPDATED: 30 Jul 2008 (20080730/ED)

Caplus now includes complete International Patent Classification (IPC) reclassification data for the second quarter of 2008.

Effective October 17, 2005, revised CAS Information Use Policies apply. They are available for your review at:

<http://www.cas.org/legal/infopolicy.html>
 'OBI' IS DEFAULT SEARCH FIELD FOR 'CAPLUS' FILE

L2 1 SEA FILE=REGISTRY ABB=ON 13981-52-7
 L3 1 SEA FILE=REGISTRY ABB=ON 14255-04-0
 L4 3024 SEA FILE=CAPLUS ABB=ON L2
 L5 4376 SEA FILE=CAPLUS ABB=ON L3
 L6 1219 SEA FILE=CAPLUS ABB=ON L4 AND L5
 L8 2364 SEA FILE=CAPLUS ABB=ON RADIATION SOURCES/CT
 L9 4 SEA FILE=CAPLUS ABB=ON L6 AND L8

L2 1 SEA FILE=REGISTRY ABB=ON 13981-52-7
 L3 1 SEA FILE=REGISTRY ABB=ON 14255-04-0
 L4 3024 SEA FILE=CAPLUS ABB=ON L2
 L5 4376 SEA FILE=CAPLUS ABB=ON L3
 L6 1219 SEA FILE=CAPLUS ABB=ON L4 AND L5
 L7 28 SEA FILE=CAPLUS ABB=ON L2/P AND L3/P
 L10 903425 SEA FILE=CAPLUS ABB=ON A/OBI
 L12 6038 SEA FILE=CAPLUS ABB=ON L10(L) (SOURCE#/OBI OR EMIT?/OBI)
 L13 28 SEA FILE=CAPLUS ABB=ON L12 AND L6
 L15 258343 SEA FILE=CAPLUS ABB=ON SEAL?/BI
 L16 2 SEA FILE=CAPLUS ABB=ON (L13 OR L7) AND L15

L2 1 SEA FILE=REGISTRY ABB=ON 13981-52-7
 L3 1 SEA FILE=REGISTRY ABB=ON 14255-04-0
 L7 28 SEA FILE=CAPLUS ABB=ON L2/P AND L3/P
 L20 248 SEA FILE=REGISTRY ABB=ON RADON?/CN
 L21 25986 SEA FILE=CAPLUS ABB=ON L20
 L25 6 SEA FILE=CAPLUS ABB=ON L7 AND L21

L2 1 SEA FILE=REGISTRY ABB=ON 13981-52-7
 L3 1 SEA FILE=REGISTRY ABB=ON 14255-04-0
 L4 3024 SEA FILE=CAPLUS ABB=ON L2
 L5 4376 SEA FILE=CAPLUS ABB=ON L3
 L6 1219 SEA FILE=CAPLUS ABB=ON L4 AND L5
 L10 903425 SEA FILE=CAPLUS ABB=ON A/OBI
 L12 6038 SEA FILE=CAPLUS ABB=ON L10(L) (SOURCE#/OBI OR EMIT?/OBI)
 L13 28 SEA FILE=CAPLUS ABB=ON L12 AND L6
 L20 248 SEA FILE=REGISTRY ABB=ON RADON?/CN
 L21 25986 SEA FILE=CAPLUS ABB=ON L20
 L24 584884 SEA FILE=CAPLUS ABB=ON 71/SC, SX ==NUCLEAR TECHNOLOGY
 L26 4 SEA FILE=CAPLUS ABB=ON L13 AND L21 AND L24

L2 1 SEA FILE=REGISTRY ABB=ON 13981-52-7
 L3 1 SEA FILE=REGISTRY ABB=ON 14255-04-0
 L4 3024 SEA FILE=CAPLUS ABB=ON L2
 L5 4376 SEA FILE=CAPLUS ABB=ON L3
 L6 1219 SEA FILE=CAPLUS ABB=ON L4 AND L5
 L7 28 SEA FILE=CAPLUS ABB=ON L2/P AND L3/P
 L10 903425 SEA FILE=CAPLUS ABB=ON A/OBI
 L12 6038 SEA FILE=CAPLUS ABB=ON L10(L) (SOURCE#/OBI OR EMIT?/OBI)
 L13 28 SEA FILE=CAPLUS ABB=ON L12 AND L6
 L18 76855 SEA FILE=CAPLUS ABB=ON POLYCARBONATE#/BI
 L19 353488 SEA FILE=CAPLUS ABB=ON HYDROXIDE#/BI

L31 1420531 SEA FILE=CAPLUS ABB=ON FILM#/BI
 L32 4 SEA FILE=CAPLUS ABB=ON (L13 OR L7) AND (L18 OR L19 OR L31)

L2 1 SEA FILE=REGISTRY ABB=ON 13981-52-7
 L3 1 SEA FILE=REGISTRY ABB=ON 14255-04-0
 L4 3024 SEA FILE=CAPLUS ABB=ON L2
 L5 4376 SEA FILE=CAPLUS ABB=ON L3
 L33 5 SEA FILE=CAPLUS ABB=ON L4(L)PUR/RL
 L34 5 SEA FILE=CAPLUS ABB=ON L5(L)PUR/RL
 L35 2 SEA FILE=CAPLUS ABB=ON L33 AND L34

=> s 19,116,125,126,132,135

L82 14 (L9 OR L16 OR L25 OR L26 OR L32 OR L35)

=> fil wpix; d que 181

FILE 'WPIX' ENTERED AT 14:09:14 ON 31 JUL 2008
 COPYRIGHT (C) 2008 THOMSON REUTERS

FILE LAST UPDATED: 29 JUL 2008 <20080729/UP>
 MOST RECENT THOMSON SCIENTIFIC UPDATE: 200848 <200848/DW>
 DERWENT WORLD PATENTS INDEX SUBSCRIBER FILE, COVERS 1963 TO DATE
 >>> Now containing more than 1.1 million chemical structures in DCR <<<

>>> IPC Reform backfile reclassifications have been loaded to the end of
 March 2008. No update date (UP) has been created for the
 reclassified documents, but they can be identified by
 20060101/UPIC and 20061231/UPIC, 20070601/UPIC, 20071001/UPIC,
 20071130/UPIC and 20080401/UPIC.
 ECLA reclassifications to April and US national classifications to
 the end of January 2008 have also been loaded. Update dates
 20080401/UPEC and /UPNC have been assigned to these. <<<

FOR A COPY OF THE DERWENT WORLD PATENTS INDEX STN USER GUIDE,
 PLEASE VISIT:

http://www.stn-international.de/training_center/patents/stn_guide.pdf

FOR DETAILS OF THE PATENTS COVERED IN CURRENT UPDATES, SEE
<http://scientific.thomsonreuters.com/support/patents/coverage/latestupdates/>

EXPLORE DERWENT WORLD PATENTS INDEX IN STN ANAVIST, VERSION 2.0:
http://www.stn-international.com/archive/presentations/DWPAnaVist2_0710.pdf

>>> HELP for European Patent Classifications see HELP ECLA, HELP ICO <<<

>>> Please note that the COPYRIGHT notification has changed <<<

'BI ABEX' IS DEFAULT SEARCH FIELD FOR 'WPIX' FILE

L70 68 SEA FILE=WPIX ABB=ON (POLONIUM/BI,ABEX OR PO/BI,ABEX) (A)210/BI
 ,ABEX OR 210PO/BI,ABEX OR PO210/BI,ABEX OR POLONIUM210/BI,ABEX
 OR 210POLONIUM/BI,ABEX
 L71 68 SEA FILE=WPIX ABB=ON (LEAD/BI,ABEX OR PB/BI,ABEX) (A)210/BI,ABE
 X OR 210PB/BI,ABEX OR PB210/BI,ABEX OR 210LEAD/BI,ABEX OR
 LEAD210/BI,ABEX

L73 974355 SEA FILE=WPIX ABB=ON FILM#/BI, ABEX
 L75 66573 SEA FILE=WPIX ABB=ON POLYCARBONATE#/BI, ABEX OR POLY CARBONATE#
 /BI, ABEX
 L76 146392 SEA FILE=WPIX ABB=ON HYDROXIDE#/BI, ABEX
 L77 722352 SEA FILE=WPIX ABB=ON SEAL?/BI, ABEX
 L78 1259 SEA FILE=WPIX ABB=ON RANDOM PULS?/BI, ABEX
 L80 2652 SEA FILE=WPIX ABB=ON ALPHA/BI, ABEX(2A) (SOURCE/BI, ABEX OR
 EMIT?/BI, ABEX OR PARTICLE#/BI, ABEX)
 L81 4 SEA FILE=WPIX ABB=ON L70 AND L71 AND (L73 OR L75 OR L76 OR
 L77 OR L78 OR L80)

=> fil PASCAL, BIOSIS, GEOREF, ENERGY, DISSABS, CONFSCI, INSPEC, EMBASE, COMPENDEX,
 SCISEARCH

FILE 'PASCAL' ENTERED AT 14:09:14 ON 31 JUL 2008
 Any reproduction or dissemination in part or in full,
 by means of any process and on any support whatsoever
 is prohibited without the prior written agreement of INIST-CNRS.
 COPYRIGHT (C) 2008 INIST-CNRS. All rights reserved.

FILE 'BIOSIS' ENTERED AT 14:09:14 ON 31 JUL 2008
 Copyright (c) 2008 The Thomson Corporation

FILE 'GEOREF' ENTERED AT 14:09:14 ON 31 JUL 2008
 COPYRIGHT (C) 2008 American Geological Institute (AGI)

FILE 'ENERGY' ENTERED AT 14:09:14 ON 31 JUL 2008
 COPYRIGHT (c) 2008 USDOE for the IEA-Energy Technology Data Exchange (ETDE)

FILE 'DISSABS' ENTERED AT 14:09:14 ON 31 JUL 2008
 COPYRIGHT (C) 2008 ProQuest Information and Learning Company; All Rights Reserved.

FILE 'CONFSCI' ENTERED AT 14:09:14 ON 31 JUL 2008
 COPYRIGHT (C) 2008 Cambridge Scientific Abstracts (CSA)

FILE 'INSPEC' ENTERED AT 14:09:14 ON 31 JUL 2008
 Compiled and produced by the IET in association WITH FIZ KARLSRUHE
 COPYRIGHT 2008 (c) THE INSTITUTION OF ENGINEERING AND TECHNOLOGY (IET)

FILE 'EMBASE' ENTERED AT 14:09:14 ON 31 JUL 2008
 Copyright (c) 2008 Elsevier B.V. All rights reserved.

FILE 'COMPENDEX' ENTERED AT 14:09:14 ON 31 JUL 2008
 Compendex Compilation and Indexing (C) 2008
 Elsevier Engineering Information Inc (EEI). All rights reserved.
 Compendex (R) is a registered Trademark of Elsevier Engineering Information Inc.

FILE 'SCISEARCH' ENTERED AT 14:09:14 ON 31 JUL 2008
 Copyright (c) 2008 The Thomson Corporation

=> d que 149; d que 154; d que 156; d que 157; d que 163; d que 168

L37 7713 SEA (POLONIUM OR PO)(A) 210 OR 210PO OR PO210
 L38 17263 SEA (LEAD OR PB)(A) 210 OR 210PB OR PB210
 L42 47252 SEA POLYCARBONATE# OR POLY CARBONATE#
 L49 2 SEA L37 AND L38 AND L42

L37 7713 SEA (POLONIUM OR PO) (A) 210 OR 210PO OR PO210
 L38 17263 SEA (LEAD OR PB) (A) 210 OR 210PB OR PB210
 L39 2217612 SEA FILM#
 L40 68820 SEA RADON OR 222RADON OR RADON222
 L41 98370 SEA ALPHA(2A) (SOURCE OR EMIT? OR PARTICLE#)
 L43 190082 SEA HYDROXIDE#
 L44 226644 SEA SEAL?
 L54 9 SEA L37 AND L38 AND L39 AND ((L40 OR L41 OR L43 OR L44))

L37 7713 SEA (POLONIUM OR PO) (A) 210 OR 210PO OR PO210
 L38 17263 SEA (LEAD OR PB) (A) 210 OR 210PB OR PB210
 L40 68820 SEA RADON OR 222RADON OR RADON222
 L41 98370 SEA ALPHA(2A) (SOURCE OR EMIT? OR PARTICLE#)
 L43 190082 SEA HYDROXIDE#
 L44 226644 SEA SEAL?
 L56 6 SEA L37 AND L38 AND L43 AND ((L40 OR L41 OR L44))

L37 7713 SEA (POLONIUM OR PO) (A) 210 OR 210PO OR PO210
 L38 17263 SEA (LEAD OR PB) (A) 210 OR 210PB OR PB210
 L40 68820 SEA RADON OR 222RADON OR RADON222
 L41 98370 SEA ALPHA(2A) (SOURCE OR EMIT? OR PARTICLE#)
 L44 226644 SEA SEAL?
 L57 4 SEA L37 AND L38 AND L44 AND ((L40 OR L41))

L37 7713 SEA (POLONIUM OR PO) (A) 210 OR 210PO OR PO210
 L38 17263 SEA (LEAD OR PB) (A) 210 OR 210PB OR PB210
 L40 68820 SEA RADON OR 222RADON OR RADON222
 L58 2389 SEA L37(2A) L38
 L61 907 SEA L40(5A) COLLECT?
 L63 1 SEA L58 AND L61

L37 7713 SEA (POLONIUM OR PO) (A) 210 OR 210PO OR PO210
 L38 17263 SEA (LEAD OR PB) (A) 210 OR 210PB OR PB210
 L40 68820 SEA RADON OR 222RADON OR RADON222
 L41 98370 SEA ALPHA(2A) (SOURCE OR EMIT? OR PARTICLE#)
 L58 2389 SEA L37(2A) L38
 L66 1616 SEA RANDOM PULS?
 L68 5 SEA L58 AND (L40 OR L41) AND L66

=> s 149,154,156,157,163,168

L83 23 (L49 OR L54 OR L56 OR L57 OR L63 OR L68)

=> dup rem 182,181,183

FILE 'CAPLUS' ENTERED AT 14:09:22 ON 31 JUL 2008
 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.
 PLEASE SEE "HELP USAGETERMS" FOR DETAILS.
 COPYRIGHT (C) 2008 AMERICAN CHEMICAL SOCIETY (ACS)

FILE 'WPIX' ENTERED AT 14:09:22 ON 31 JUL 2008

COPYRIGHT (C) 2008 THOMSON REUTERS

FILE 'BIOSIS' ENTERED AT 14:09:22 ON 31 JUL 2008
Copyright (c) 2008 The Thomson CorporationFILE 'ENERGY' ENTERED AT 14:09:22 ON 31 JUL 2008
COPYRIGHT (c) 2008 USDOE for the IEA-Energy Technology Data Exchange (ETDE)FILE 'INSPEC' ENTERED AT 14:09:22 ON 31 JUL 2008
Compiled and produced by the IET in association WITH FIZ KARLSRUHE
COPYRIGHT 2008 (c) THE INSTITUTION OF ENGINEERING AND TECHNOLOGY (IET)FILE 'EMBASE' ENTERED AT 14:09:22 ON 31 JUL 2008
Copyright (c) 2008 Elsevier B.V. All rights reserved.FILE 'COMPENDEX' ENTERED AT 14:09:22 ON 31 JUL 2008
Compendex Compilation and Indexing (C) 2008
Elsevier Engineering Information Inc (EEI). All rights reserved.
Compendex (R) is a registered Trademark of Elsevier Engineering Information Inc.FILE 'SCISEARCH' ENTERED AT 14:09:22 ON 31 JUL 2008
Copyright (c) 2008 The Thomson Corporation
PROCESSING COMPLETED FOR L82
PROCESSING COMPLETED FOR L81
PROCESSING COMPLETED FOR L83
L84 31 DUP REM L82 L81 L83 (10 DUPLICATES REMOVED)
ANSWERS '1-14' FROM FILE CAPLUS
ANSWERS '15-16' FROM FILE WPIX
ANSWERS '17-20' FROM FILE BIOSIS
ANSWERS '21-28' FROM FILE ENERGY
ANSWER '29' FROM FILE EMBASE
ANSWER '30' FROM FILE COMPENDEX
ANSWER '31' FROM FILE SCISEARCH

=> d ibib ab hitind 1-14; d iall abex tech 15-16; d iall 17-31; fil hom

L84 ANSWER 1 OF 31 CAPLUS COPYRIGHT 2008 ACS on STN DUPLICATE 1
ACCESSION NUMBER: 2004:1156703 CAPLUS Full-text
DOCUMENT NUMBER: 142:44783
TITLE: Method for producing a sealed 210Pb-210Po
alpha source (alpha
particle emitter) and apparatus thereof
INVENTOR(S): Mitsugashira, Hiroaki; Tsuyuzaki, Noriyoshi
PATENT ASSIGNEE(S): Japan
SOURCE: PCT Int. Appl., 21 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

| PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|--|------|----------|-----------------|----------|
| WO 2004114324 | A2 | 20041229 | WO 2004-JP8407 | 20040609 |
| WO 2004114324 | A3 | 20050224 | | |
| W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, | | | | |

TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
 RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM,
 AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK,
 EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE,
 SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE,
 SN, TD, TG

| | | | | |
|--|----|----------|------------------|----------|
| JP 2005010009 | A | 20050113 | JP 2003-174296 | 20030619 |
| EP 1634302 | A2 | 20060315 | EP 2004-736467 | 20040609 |
| R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, FI, RO, CY, TR, BG, CZ, EE, HU, PL, SK | | | | |
| CN 1826663 | A | 20060830 | CN 2004-80017060 | 20040609 |
| US 20070098606 | A1 | 20070503 | US 2005-560922 | 20051215 |
| JP 2003-174296 A 20030619 | | | | |
| WO 2004-JP8407 W 20040609 | | | | |

PRIORITY APPLN. INFO.:

AB A method is described for producing a sealed 210Pb-210Po α source for a random pulse generator. The method includes: collecting 210Pb-210Po with a 210Pb collector using Rn collection; precipitating the hydroxides of the collected 210Pb-210Po and collecting the ppt. by a polycarbonate filter; dissolving the 210Pb-210Po hydroxide precipitate to form a 210Pb-210Po radioactive thin film ; and sealing the 210Pb-210Po radioactive thin film for protection.

IC ICM G21G004-00

CC 71-6 (Nuclear Technology)

ST radiation source lead polonium radon alpha particle emitter

IT Radiation sources
 (method for producing a sealed 210Pb-210Po alpha source)

IT Polycarbonates, uses
 Uranium ores
 RL: NUU (Other use, unclassified); USES (Uses)
 (method for producing a sealed 210Pb-210Po alpha source)

IT 13981-52-7P, Polonium-210, preparation 14255-04-0P,
 Lead-210, preparation
 RL: PUR (Purification or recovery); PREP (Preparation)
 (method for producing a sealed 210Pb-210Po alpha source)

IT 14859-67-7, Radon-222, uses
 RL: TEM (Technical or engineered material use); USES (Uses)
 (method for producing a sealed 210Pb-210Po alpha source)

L84 ANSWER 2 OF 31 CAPLUS COPYRIGHT 2008 ACS on STN DUPLICATE 2

ACCESSION NUMBER: 2003:91799 CAPLUS Full-text

DOCUMENT NUMBER: 138:261770

TITLE: Collection of emanating 222Rn for the preparation of a 210Pb-210Po alpha-source and the building of a mobile random pulse and probability generator utilizing alpha-counting technique

AUTHOR(S): Hirose, N.; Tsuyuzaki, N.; Yamamot, H.; Mitsugashira, T.; Hara, M.

CORPORATE SOURCE: IWAKI Electronics Co., Ltd., Iwaki-city, Fukushima, 972-8322, Japan

SOURCE: Journal of Radioanalytical and Nuclear Chemistry (2003), 255(1), 207-210

CODEN: JRNCDM; ISSN: 0236-5731

PUBLISHER: Kluwer Academic Publishers

DOCUMENT TYPE: Journal

LANGUAGE: English

AB A random pulse and probability generator (RPG) was developed using the detection technique of alpha-particles as the random signal source. The collection technique for 222Rn emanated from natural U ore was examined for preparing highly pure 210Pb-210Po as an alpha source for RPG. The yield with a trap refrigerated by liquid N is >99% for 222Rn collection.

CC 71-7 (Nuclear Technology)

ST alpha source probability generator

IT Distribution function
(Poisson; in relation to alpha-source for random pulse and probability generator)

IT Probability
(emanating 222Rn in preparation of 210Pb-210Po alpha-source for random pulse and probability generator)

IT 12587-46-1, Alpha particle
RL: PEP (Physical, engineering or chemical process); PYP (Physical process); PROC (Process)
(decay detector; alpha-source for random pulse and probability generator)

IT 14859-67-7, Radon-222, uses
RL: NUU (Other use, unclassified); USES (Uses)
(emanating 222Rn in preparation of 210Pb-210Po alpha-source for random pulse and probability generator)

IT 13981-52-7, Polonium-210, processes 14255-04-0,
Lead-210, processes
RL: PEP (Physical, engineering or chemical process); PYP (Physical process); PROC (Process)
(emanating 222Rn in preparation of 210Pb-210Po alpha-source for random pulse and probability generator)

REFERENCE COUNT: 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L84 ANSWER 3 OF 31 CAPLUS COPYRIGHT 2008 ACS on STN DUPLICATE 3
ACCESSION NUMBER: 2000:551944 CAPLUS Full-text
DOCUMENT NUMBER: 133:156747
TITLE: Radiation sources for examination and/or calibration of radiation detectors
INVENTOR(S): Von Philipsborn, Henning
PATENT ASSIGNEE(S): Germany
SOURCE: Ger. Offen., 8 pp.
CODEN: GWXXBX
DOCUMENT TYPE: Patent
LANGUAGE: German
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

| PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|------------------------|------|----------|------------------|-------------|
| DE 10002113 | A1 | 20000810 | DE 2000-10002113 | 20000119 |
| PRIORITY APPLN. INFO.: | | | DE 1999-19902145 | A1 19990120 |

AB A source for calibration comprises a radioelement-enriched carrier substrate. The radionuclide may be Pb-210 and/or Po-210. 1St a Rn-222-free substance is brought into a gas volume and then the Rn-222 concentration in the volume is increased. Then the gas contained in the volume is introduced into an organic solvent, whereby the Rn-222 contained in the gas dissolves in the solvent. The solvent is then aged over several days and poured over the carrier substrate through a screen end glass-fiber filter. Alternatively the carrier substrate can be also temporarily stored in the solvent. The process is low-cost and safe.

IC ICM G21G004-04
ICS G01T001-16

CC 71-6 (Nuclear Technology)
 IT Calibration
 Radiation detectors
 Radiation sources
 (radiation sources for examination and/or calibration of radiation detectors)
 IT 13981-52-7, Polonium-210, processes 14255-04-0,
 Lead-210, processes 14859-67-7, Radon-222, processes
 RL: PEP (Physical, engineering or chemical process); PROC (Process)
 (radiation sources for examination and/or calibration of radiation detectors)

L84 ANSWER 4 OF 31 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 2007:71900 CAPLUS Full-text
 DOCUMENT NUMBER: 147:352120
 TITLE: Estimate computing on source of nuclear and radiological terrorism events
 AUTHOR(S): Fu, Guang-zhi; Liu, Jun-feng; He, Bin; Zhang, Xi-xi
 CORPORATE SOURCE: The Second Artillery Engineering Institute, Xian of Shaanxi Prov., 710025, Peop. Rep. China
 SOURCE: He Dianzixue Yu Tance Jishu (2006), 26(6), 723-725, 716
 CODEN: HDYUEC; ISSN: 0258-0934
 PUBLISHER: Yuanzinen Chubanshe
 DOCUMENT TYPE: Journal
 LANGUAGE: Chinese

AB Once the nuclear and radiol. terrorism events have happened, and we do not know the nuclear kinds and amts., it should be very difficult for us to make a reasonable appraisement about the events, at the same time it might affect the protective and active, decision-making about meeting the emergency. One source anal. method was presented in this paper, and this method was used to explain an example of the source about the nuclear and radiol. events. This paper also pointed out that the mainly radiol. dangerous in the nuclear and radiol. terrorism events was long-lived α aerosol, and the mainly chemical dangerous was the very poisonous nuclide.

CC 71-14 (Nuclear Technology)
 ST source nuclear radiol terrorism event alpha aerosol nuclide
 IT 7440-69-9, Bismuth (209), formation (nonpreparative) 13966-01-3, Thallium (210), formation (nonpreparative) 13966-27-3, Lead (206), formation (nonpreparative) 13966-29-5, Uranium (234), formation (nonpreparative) 13968-55-3, Uranium (233), formation (nonpreparative) 13981-14-1, Protactinium (233), formation (nonpreparative) 13981-52-7, Polonium (210), formation (nonpreparative) 13981-53-8, Radium (225), formation (nonpreparative) 13982-10-0, Plutonium (242), formation (nonpreparative) 13982-63-3, Radium (226), formation (nonpreparative) 13994-20-2, Neptunium (237), formation (nonpreparative) 14119-29-0, Lead (207), formation (nonpreparative) 14119-30-3, Lead (209), formation (nonpreparative) 14119-32-5, Plutonium (241), formation (nonpreparative) 14119-33-6, Plutonium (240), formation (nonpreparative) 14133-67-6, Thallium (207), formation (nonpreparative) 14255-04-0, Lead (210), formation (nonpreparative) 14265-85-1, Actinium (225), formation (nonpreparative) 14269-63-7, Thorium (230), formation (nonpreparative) 14331-79-4, Bismuth (210), formation (nonpreparative) 14331-85-2, Protactinium (231), formation (nonpreparative) 14596-10-2, Americium (241), formation (nonpreparative) 14733-03-0, Bismuth (214), formation (nonpreparative) 14835-02-0, Radon (219), formation (nonpreparative) 14859-67-7, Radon (222), formation (nonpreparative) 14932-40-2, Thorium (231), formation (nonpreparative) 15035-09-3, Thallium (206), formation (nonpreparative)

15065-10-8, Thorium (234), formation (nonpreparative) 15067-28-4, Lead (214), formation (nonpreparative) 15100-28-4, Protactinium (234), formation (nonpreparative) 15117-48-3, Plutonium (239), formation (nonpreparative) 15117-96-1, Uranium (235), formation (nonpreparative) 15229-37-5, Bismuth (211), formation (nonpreparative) 15422-74-9, Polonium (218), formation (nonpreparative) 15594-54-4, Thorium (229), formation (nonpreparative) 15623-45-7, Radium (223), formation (nonpreparative) 15623-47-9, Thorium (227), formation (nonpreparative) 15690-73-0, Thallium (209), formation (nonpreparative) 15706-52-2, Polonium (215), formation (nonpreparative) 15735-67-8, Polonium (214), formation (nonpreparative) 15735-83-8, Polonium (211), formation (nonpreparative) 15755-40-5, Astatine (218), formation (nonpreparative) 15756-41-9, Francium (221), formation (nonpreparative) 15756-57-7, Polonium (213), formation (nonpreparative) 15756-98-6, Francium (223), formation (nonpreparative) 15776-20-2, Bismuth (213), formation (nonpreparative) 15816-77-0, Lead (211), formation (nonpreparative) 17239-90-6, Astatine (217), formation (nonpreparative) 51696-22-1, Actinium (237), formation (nonpreparative) 51696-49-2, Thorium (237), formation (nonpreparative)
 RL: FMU (Formation, unclassified); POL (Pollutant); FORM (Formation, nonpreparative); OCCU (Occurrence)
 (estimate computing on source of nuclear and radiol. terrorism events)

L84 ANSWER 5 OF 31 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2006:256884 CAPLUS Full-text

DOCUMENT NUMBER: 144:365398

TITLE: Performance characteristics of sequential separation and quantification of lead-210 and polonium-210 by ion exchange chromatography and nuclear spectrometric measurements

AUTHOR(S): El Afifi, E. M.; Borai, E. H.

CORPORATE SOURCE: Hot Laboratories and Waste Management Center (HLWMC), Atomic Energy Authority, Cairo, Egypt

SOURCE: Journal of Environmental Quality (2006), 35(2), 568-574

CODEN: JEVQAA; ISSN: 0047-2425

PUBLISHER: American Society of Agronomy

DOCUMENT TYPE: Journal

LANGUAGE: English

AB A selective separation and quant. determination procedure for 210Pb and 210Po in various environmental matrixes from different sources such as IAEA-326 soil, phosphate rocks (PR), and phosphogypsum (PG) was developed. The tested samples were digested sequentially using concentrated mineral acids (HF, HNO3) by a programmable high-pressure microwave digestion system. The sample solution was loaded onto a preconditioned ion exchange column (Sr-resin) for chromatog. separation Polonium-210 was eluted by 6 M HNO3 then spontaneously deposited onto polished silver disks to be measured using low-background alpha spectrometry. Lead-210 was sequentially eluted using 6 M HCl solution, precipitated as lead oxalate, dissolved in HNO3 solution, and mixed with scintillation cocktail to be measured by liquid scintillation counting (LSC). Performance of the developed procedure was tested using a reference soil (IAEA-326), with recommended isotope values, that was used as a quality control to assess separation and quantification efficiency (recovery %). The min. detectable activities of 210Pb and 210Po were found to be 24 and 0.28 Bq kg-1 for the measurements using LSC and alpha spectrometry, resp. The recoveries (%) of 210Pb and 210Po were found to be 80 and 60%, resp. To test the validity of the proposed LSC method, a comparative study was performed by measuring 210Pb activity concentration in test samples by nondestructive gamma-ray spectrometry.

CC 8-1 (Radiation Biochemistry)

Section cross-reference(s): 19
 IT 13981-52-7P, Polonium-210, analysis 14255-04-0P,
 Lead-210, analysis
 RL: ANT (Analyte); PUR (Purification or recovery); ANST
 (Analytical study); PREP (Preparation)
 (sequential separation and quantification of lead-210 and polonium-210 by
 ion exchange chromatog. and nuclear spectrometric measurements)
 REFERENCE COUNT: 31 THERE ARE 31 CITED REFERENCES AVAILABLE FOR THIS
 RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L84 ANSWER 6 OF 31 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 2003:535419 CAPLUS Full-text
 DOCUMENT NUMBER: 139:313232
 TITLE: Gamma coincidence study of $^{208}\text{Pb} + ^{350}\text{MeV } ^{64}\text{Ni}$
 collisions
 AUTHOR(S): Krolas, W.; Broda, R.; Fornal, B.; Pawlat, T.; Grawe,
 H.; Maier, K. H.; Schramm, M.; Schubart, R.
 CORPORATE SOURCE: H. Niewodniczanski Institute of Nuclear Physics,
 Krakow, PL-31342, Pol.
 SOURCE: Nuclear Physics A (2003), A724(3,4), 289-312
 CODEN: NUPABL; ISSN: 0375-9474
 PUBLISHER: Elsevier Science B.V.
 DOCUMENT TYPE: Journal
 LANGUAGE: English
 AB Products of $^{208}\text{Pb} + ^{64}\text{Ni}$ collisions at an energy 12% above the Coulomb barrier
 were studied in a γ -spectroscopy thick target experiment. The product yield
 distribution was established from the γ - γ coincidence anal. supplemented by
 target radioactivity measurements. Neutron evaporation from excited primary
 products was estimated to determine the pre-emission map of fragments. We
 discuss the transfer of protons and neutrons between the colliding ions in
 terms of the N/Z ratio equilibration.

CC 70-1 (Nuclear Phenomena)
 IT 7439-96-5P, Manganese 55, preparation 7440-03-1P, Niobium 93,
 preparation 7440-25-7P, Tantalum 181, preparation 7440-38-2P, Arsenic
 75, preparation 7440-48-4P, Cobalt 59, preparation 7440-57-5P, Gold
 197, preparation 7440-65-5P, Yttrium 89, preparation 7440-69-9P,
 Bismuth 209, preparation 10043-49-9P, Gold 198, preparation
 10098-91-6P, Yttrium 90, preparation 10098-97-2P, Strontium 90,
 preparation 10198-40-0P, Cobalt 60, preparation 13965-98-5P, Krypton
 83, preparation 13966-06-8P, Tin 113, preparation 13966-26-2P, Lead
 204, preparation 13966-27-3P, Lead 206, preparation 13966-31-9P,
 Manganese 54, preparation 13967-66-3P, Iridium 191, preparation
 13967-67-4P, Iridium 193, preparation 13967-71-0P, Zirconium 95,
 preparation 13967-73-2P, Strontium 85, preparation 13967-74-3P, Cerium
 141, preparation 13967-76-5P, Niobium 95, preparation 13968-47-3P,
 Iron 58, preparation 13968-51-9P, Thallium 204, preparation
 13968-53-1P, Ruthenium 103, preparation 13981-21-0P, Mercury 198,
 preparation 13981-25-4P, Copper 64, preparation 13981-27-6P, Zirconium
 89, preparation 13981-29-8P, Terbium 160, preparation 13981-32-3P,
 Selenium 76, preparation 13981-37-8P, Nickel 63, preparation
 13981-38-9P, Cobalt 58, preparation 13981-50-5P, Cobalt 57, preparation
 13981-51-6P, Mercury 197, preparation 13981-52-7P, Polonium 210,
 preparation 13981-59-4P, Tin 117, preparation 13981-78-7P, Chromium
 53, preparation 13981-80-1P, Nickel 60, preparation 13981-81-2P,
 Nickel 62, preparation 13981-83-4P, Cobalt 61, preparation
 13981-84-5P, Cobalt 63, preparation 13982-00-8P, Tantalum 182,
 preparation 13982-08-6P, Ytterbium 170, preparation 13982-09-7P,
 Osmium 186, preparation 13982-12-2P, Rubidium 85, preparation
 13982-13-3P, Rubidium 87, preparation 13982-14-4P, Strontium 86,
 preparation 13982-15-5P, Zirconium 90, preparation 13982-20-2P, Gold

193, preparation 13982-21-3P, Germanium 72, preparation 13982-22-4P, Gallium 72, preparation 13982-23-5P, Zinc 69, preparation 13982-24-6P, Iron 54, preparation 13982-30-4P, Cerium 139, preparation 13982-36-0P, Yttrium 88, preparation 13982-37-1P, Niobium 92, preparation 13982-38-2P, Bismuth 207, preparation 13982-39-3P, Zinc 65, preparation 13982-64-4P, Strontium 87, preparation 13982-78-0P, Mercury 203, preparation 13983-27-2P, Krypton 85, preparation 13994-19-9P, Xenon 127, preparation 14041-45-3P, Ytterbium 167, preparation 14041-50-0P, Ytterbium 171, preparation 14041-51-1P, Ytterbium 173, preparation 14041-52-2P, Ytterbium 172, preparation 14041-58-8P, Cadmium 114, preparation 14092-98-9P, Chromium 52, preparation 14093-02-8P, Iron 56, preparation 14093-09-5P, Hafnium 177, preparation 14093-11-9P, Hafnium 172, preparation 14093-12-0P, Lutetium 172, preparation 14107-52-9P, Thallium 197, preparation 14119-06-3P, Copper 65, preparation 14119-10-9P, Strontium 88, preparation 14119-12-1P, Zirconium 94, preparation 14119-15-4P, Molybdenum 99, preparation 14119-17-6P, Tin 120, preparation 14119-18-7P, Tin 122, preparation 14119-24-5P, Osmium 191, preparation 14119-28-9P, Lead 205, preparation 14119-29-0P, Lead 207, preparation 14119-30-3P, Lead 209, preparation 14133-67-6P, Thallium 207, preparation 14133-76-7P, Technetium 99, preparation 14145-42-7P, Bismuth 208, preparation 14158-27-1P, Strontium 89, preparation 14158-30-6P, Iodine 124, preparation 14158-35-1P, Iridium 194, preparation 14191-65-2P, Rubidium 89, preparation 14191-69-6P, Indium 116, preparation 14191-70-9P, Tin 116, preparation 14191-81-2P, Krypton 82, preparation 14191-82-3P, Krypton 86, preparation 14191-84-5P, Copper 63, preparation 14191-86-7P, Mercury 202, preparation 14191-87-8P, Mercury 199, preparation 14191-88-9P, Platinum 195, preparation 14234-24-3P, Yttrium 91, preparation 14255-04-0P, Lead 210, preparation 14265-71-5P, Selenium 75, preparation 14265-76-0P, Hafnium 179, preparation 14265-77-1P, Hafnium 178, preparation 14265-78-2P, Hafnium 180, preparation 14265-79-3P, Tungsten 180, preparation 14265-80-6P, Tungsten 182, preparation 14265-81-7P, Tungsten 183, preparation 14265-82-8P, Tungsten 184, preparation 14265-83-9P, Tungsten 186, preparation 14265-84-0P, Iridium 189, preparation 14269-78-4P, Ytterbium 169, preparation 14274-68-1P, Yttrium 87, preparation 14274-76-1P, Molybdenum 96, preparation 14274-79-4P, Osmium 190, preparation 14274-81-8P, Osmium 188, preparation 14276-53-0P, Copper 62, preparation 14280-37-6P, Bismuth 199, preparation 14280-38-7P, Bismuth 201, preparation 14280-48-9P, Thallium 203, preparation 14280-49-0P, Thallium 205, preparation 14304-78-0P, Arsenic 74, preparation 14304-79-1P, Tellurium 121, preparation 14304-80-4P, Tellurium 123, preparation 14304-97-3P, Chromium 54, preparation 14320-93-5P, Gold 195, preparation 14331-79-4P, Bismuth 210, preparation 14331-81-8P, Mercury 206, preparation 14331-90-9P, Bromine 84, preparation 14331-91-0P, Strontium 91, preparation 14331-93-2P, Zirconium 91, preparation 14333-38-1P, Bismuth 205, preparation 14336-70-0P, Nickel 59, preparation 14374-79-9P, Antimony 122, preparation 14374-81-3P, Germanium 71, preparation 14378-26-8P, Rhenium 188, preparation 14378-27-9P, Rhenium 190, preparation 14378-32-6P, Zinc 64, preparation 14378-33-7P, Zinc 66, preparation 14378-34-8P, Zinc 67, preparation 14378-35-9P, Zinc 68, preparation 14378-36-0P, Zinc 70, preparation 14378-53-1P, Rhodium 101, preparation 14380-59-7P, Bromine 81, preparation 14391-02-7P, Gallium 69, preparation 14391-03-8P, Gallium 71, preparation 14391-10-7P, Terbium 156, preparation 14391-11-8P, Gold 199, preparation 14391-27-6P, Tantalum 179, preparation 14391-28-7P, Rhenium 185, preparation 14391-29-8P, Rhenium 187, preparation 14391-61-8P, Bromine 80, preparation 14391-63-0P, Rubidium 82, preparation 14391-68-5P, Antimony 120, preparation 14391-73-2P, Copper 66, preparation

14391-74-3P, Gallium 70, preparation 14391-76-5P, Silver 110, preparation 14392-07-5P, Gadolinium 156, preparation 14392-15-5P, Zirconium 92, preparation 14392-17-7P, Molybdenum 95, preparation 14392-19-9P, Molybdenum 97, preparation 14392-20-2P, Molybdenum 98, preparation 14392-21-3P, Molybdenum 100, preparation 14452-48-3P, Hafnium 176, preparation 14484-13-0P, Rhenium 183, preparation 14596-12-4P, Iron 59, preparation 14681-52-8P, Manganese 56, preparation 14681-54-0P, Selenium 80, preparation 14681-59-5P, Iron 55, preparation 14681-63-1P, Niobium 94, preparation 14681-65-3P, Niobium 90, preparation 14681-72-2P, Selenium 77, preparation 14682-97-4P, Niobium 91, preparation 14683-00-2P, Molybdenum 94, preparation 14683-10-4P, Antimony 124, preparation 14683-19-3P, Promethium 148, preparation 14683-24-0P, Gadolinium 154, preparation 14683-25-1P, Dysprosium 160, preparation 14683-29-5P, Ytterbium 174, preparation 14683-32-0P, Tungsten 179, preparation 14683-36-4P, Tantalum 183, preparation 14686-69-2P, Bromine 82, preparation 14687-25-3P, Lead 203, preparation 14687-40-2P, Germanium 75, preparation 14687-41-3P, Germanium 76, preparation 14687-50-4P, Bismuth 202, preparation 14687-58-2P, Selenium 82, preparation 14687-59-3P, Germanium 77, preparation 14687-60-6P, Selenium 83, preparation 14687-61-7P, Arsenic 77, preparation 14687-62-8P, Bromine 83, preparation 14694-69-0P, Iridium 192, preparation 14762-69-7P, Iron 57, preparation 14809-52-0P, Yttrium 85, preparation 14809-53-1P, Yttrium 86, preparation 14809-60-0P, Chromium 55, preparation 14809-62-2P, Cobalt 62, preparation 14809-64-4P, Gallium 74, preparation 14809-66-6P, Arsenic 79, preparation 14809-68-8P, Krypton 87, preparation 14833-10-4P, Vanadium 53, preparation 14833-16-0P, Selenium 78, preparation 14833-43-3P, Erbium 168, preparation 14833-49-9P, Nickel 65, preparation 14834-71-0P, Iridium 187, preparation 14834-72-1P, Promethium 143, preparation 14834-83-4P, Ytterbium 166, preparation 14834-85-6P, Dysprosium 162, preparation 14867-61-9P, Platinum 196, preparation 14900-10-8P, Erbium 164, preparation 14900-11-9P, Erbium 166, preparation 14900-13-1P, Thulium 168, preparation 14900-21-1P, Hafnium 181, preparation 14903-04-9P, Bismuth 204, preparation 14913-25-8P, Dysprosium 158, preparation 14913-50-9P, Thallium 208, preparation 14913-85-0P, Platinum 192, preparation 14913-89-4P, preparation 14914-16-0P, Gold 196, preparation 14914-52-4P, Zinc 71, preparation 14914-59-1P, Palladium 106, preparation 14914-60-4P, Ruthenium 100, preparation 14914-61-5P, Ruthenium 101, preparation 14914-62-6P, Ruthenium 102, preparation 14914-65-9P, Tin 118, preparation 14914-66-0P, Indium 117, preparation 14917-67-0P, Mercury 196, preparation 14922-49-7P, Hafnium 174, preparation 14922-68-0P, Osmium 184, preparation 14922-70-4P, Platinum 188, preparation 14928-10-0P, Nickel 61, preparation 14928-36-0P, Rubidium 88, preparation 14932-41-3P, Tungsten 185, preparation 14932-53-7P, Rubidium 86, preparation 14981-91-0P, Iridium 190, preparation 14983-46-1P, Rhenium 184, preparation 14983-48-3P, Tungsten 187, preparation 14993-36-3P, Osmium 182, preparation

RL: PNU (Preparation, unclassified); PREP (Preparation)

(gamma coincidence study of 208Pb+350 MeV 64Ni collisions with light to heavy mass products)

IT 14993-62-5P, Rhenium 180, preparation 14993-65-8P, Rhenium 181, preparation 14993-91-0P, Krypton 84, preparation 14995-61-0P, Krypton 88, preparation 14998-63-1P, Rhenium 186, preparation 14998-72-2P, Tin 114, preparation 14998-96-0P, Platinum 194, preparation 14999-33-8P, Manganese 53, preparation 15034-51-2P, Gallium 73, preparation 15034-58-9P, Germanium 73, preparation 15034-59-0P, Germanium 74, preparation 15035-09-3P, Thallium 206, preparation 15047-33-3P, Rhenium 179, preparation 15055-22-8P, Tantalum 178, preparation 15055-23-9P, Tungsten 178, preparation 15055-30-8P, Platinum 189,

preparation 15062-08-5P, Osmium 192, preparation 15064-65-0P, Thallium 201, preparation 15064-66-1P, Thallium 199, preparation 15064-97-8P, Mercury 194, preparation 15116-82-2P, Mercury 193, preparation 15125-00-5P, Rubidium 90, preparation 15125-53-8P, Tin 112, preparation 15185-19-0P, Mercury 201, preparation 15229-37-5P, Bismuth 211, preparation 15389-34-1P, Polonium 212, preparation 15411-62-8P, Ruthenium 99, preparation 15422-54-5P, Iron 61, preparation 15422-58-9P, Selenium 81, preparation 15422-59-0P, Arsenic 73, preparation 15575-20-9P, Arsenic 76, preparation 15678-91-8P, Krypton 81, preparation 15690-73-0P, Thallium 209, preparation 15691-06-2P, Zirconium 96, preparation 15700-41-1P, Niobium 98, preparation 15700-42-2P, Niobium 100, preparation 15700-83-1P, Bromine 85, preparation 15701-21-0P, Tantalum 184, preparation 15706-36-2P, Platinum 191, preparation 15706-38-4P, Polonium 204, preparation 15715-01-2P, Astatine 213, preparation 15715-06-7P, Mercury 192, preparation 15720-36-2P, Cobalt 64, preparation 15720-38-4P, Copper 68, preparation 15720-40-8P, Copper 69, preparation 15720-45-3P, Polonium 207, preparation 15720-55-5P, Thallium 200, preparation 15720-57-7P, Thallium 202, preparation 15735-68-9P, Platinum 190, preparation 15735-70-3P, Platinum 193, preparation 15735-74-7P, Platinum 197, preparation 15735-81-6P, Polonium 209, preparation 15735-83-8P, Polonium 211, preparation 15735-86-1P, Polonium 206, preparation 15735-87-2P, Polonium 208, preparation 15741-32-9P, Lutetium 170, preparation 15741-33-0P, Manganese 57, preparation 15743-50-7P, Thallium 198, preparation 15743-54-1P, Ytterbium 168, preparation 15743-55-2P, Zinc 72, preparation 15749-40-3P, Titanium 52, preparation 15749-46-9P, Tungsten 181, preparation 15749-58-3P, Palladium 108, preparation 15750-13-7P, Hafnium 175, preparation 15751-77-6P, Zirconium 93, preparation 15752-22-4P, Iridium 188, preparation 15752-27-9P, Lutetium 171, preparation 15752-86-0P, Lead 202, preparation 15755-35-8P, Arsenic 78, preparation 15755-36-9P, Astatine 212, preparation 15755-38-1P, Astatine 209, preparation 15755-39-2P, Astatine 211, preparation 15756-10-2P, Mercury 200, preparation 15756-14-6P, Mercury 204, preparation 15756-15-7P, Mercury 195, preparation 15756-45-3P, Gold 192, preparation 15756-63-5P, Platinum 198, preparation 15756-69-1P, Polonium 202, preparation 15756-83-9P, Germanium 78, preparation 15756-89-5P, Gold 194, preparation 15756-97-5P, Francium 212, preparation 15757-14-9P, Gallium 68, preparation 15757-23-0P, Hafnium 173, preparation 15757-86-5P, Copper 67, preparation 15758-18-6P, Chromium 56, preparation 15758-45-9P, Selenium 79, preparation 15758-49-3P, Strontium 84, preparation 15759-29-2P, Tantalum 180, preparation 15761-06-5P, Osmium 189, preparation 15765-69-2P, Radon 211, preparation 15765-70-3P, Radon 212, preparation 15765-78-3P, Rhenium 189, preparation 15765-86-3P, Rubidium 84, preparation 15766-01-5P, Ruthenium 104, preparation 15766-16-2P, Nickel 67, preparation 15766-33-3P, Nickel 66, preparation 15766-50-4P, Osmium 185, preparation 15766-52-6P, Osmium 187, preparation 15776-19-9P, Bismuth 206, preparation 15816-77-0P, Lead 211, preparation 15816-99-6P, Iridium 195, preparation 15832-32-3P, Niobium 96, preparation 15832-38-9P, Gold 190, preparation 15832-41-4P, Indium 118, preparation 15840-05-8P, Erbium 162, preparation 16394-57-3P, Gold 191, preparation 16468-57-8P, Germanium 79, preparation 16645-99-1P, Lead 200, preparation 16646-00-7P, Lead 198, preparation 16729-68-3P, Mercury 205, preparation 16729-74-1P, Polonium 203, preparation 16729-76-3P, Polonium 205, preparation 17056-36-9P, Rubidium 83, preparation 17058-33-2P, Radon 209, preparation 17239-85-9P, Bismuth 200, preparation 17239-87-1P, Lead 201, preparation 17620-09-6P, Francium 213, preparation 17638-03-8P, Radium 214, preparation 18390-45-9P, Astatine 208, preparation 18476-92-1P,

Manganese 58, preparation 18496-04-3P, Niobium 97, preparation 18624-12-9P, Cobalt 66, preparation 18724-77-1P, Thallium 196, preparation 18830-37-0P, Astatine 210, preparation 20091-45-6P, Gold 200, preparation 20601-76-7P, Astatine 207, preparation 21402-14-2P, Gallium 75, preparation 21410-52-6P, Gallium 76, preparation 21459-51-8P, Astatine 206, preparation 21459-71-2P, Rhenium 182, preparation 21459-72-3P, Osmium 183, preparation 22453-47-0P, Platinum 187, preparation 22453-70-9P, Niobium 99, preparation 24383-94-6P, Bismuth 203, preparation 24447-13-0P, Iridium 186, preparation 25731-76-4P, Yttrium 84, preparation 26110-67-8P, Krypton 80, preparation 26683-69-2P, Thallium 195, preparation 27485-99-0P, Lead 197, preparation 27486-00-6P, Lead 199, preparation 27742-26-3P, Iridium 184, preparation 28637-43-6P, Vanadium 54, preparation 29054-43-1P, Iridium 185, preparation 29136-29-6P, Radon 214, preparation 29675-20-5P, Iron 62, preparation 29675-21-6P, Nickel 68, preparation 29675-28-3P, Copper 70, preparation 29675-32-9P, Titanium 53, preparation 29675-34-1P, Nickel 69, preparation 29675-35-2P, Zinc 73, preparation 29675-38-5P, Cobalt 65, preparation 30017-28-8P, Copper 71, preparation 32020-21-6P, Iron 60, preparation 32025-57-3P, Radon 213, preparation 33233-20-4P, Radon 210, preparation 33690-55-0P, Arsenic 80, preparation 36819-19-9P, Vanadium 55, preparation 36819-21-3P, Chromium 57, preparation 36819-22-4P, Manganese 59, preparation 42250-70-4P, Chromium 58, preparation 42250-73-7P, Manganese 60, preparation 52813-79-3P, Manganese 61, preparation 58831-77-9P, Iron 64, preparation 58831-78-0P, Iron 63, preparation 72062-02-3P, Manganese 62, preparation
RL: PNU (Preparation, unclassified); PREP (Preparation)
(gamma coincidence study of 208Pb+350 MeV 64Ni collisions with light to heavy mass products)

REFERENCE COUNT: 23 THERE ARE 23 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L84 ANSWER 7 OF 31 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2004:56675 CAPLUS Full-text
DOCUMENT NUMBER: 141:29931
TITLE: Preparation of low level sealed 210Pb source for random pulse generator
AUTHOR(S): Mitsugashira, T.; Hara, M.; Tsuyuzaki, N.
CORPORATE SOURCE: Institute for Materials Research, Tohoku University, Japan
SOURCE: KEK Proceedings (2003), 2003-11(Proceedings of the Fourth Workshop on Environmental Radioactivity, 2003), 263-268
CODEN: KEPREW
PUBLISHER: High Energy Accelerator Research Organization
DOCUMENT TYPE: Journal
LANGUAGE: Japanese

AB Two methods for sealed 210Pb-210Po source preparation, direct deposition from isoPr alc. solution (IPA solution) of 210Pb-210Po nitrate (D-IPA method) and the 210Pb-210Po hydroxides precipitation (PPT method), were exptl. examined. In the former D-IPA method, an aliquot of IPA solution of 210Pb-210Po nitrate was directly dropped in a sealed cap for PPD and dried by heating. Then, a polycarbonate (PC) solution of 1/1 mixture of CH₂Cl₂ and dichloroethane was dropped on the source to make a thin (.apprx.0.1 mg/cm²) film for radioactivity sealed. In the PPT method, 210Pb-210Po hydroxide was filtered on a PC membrane filter (Nuclipore 0.1 μ m) and the membrane filter was dissolved in a 1/1 mixture of CH₂Cl₂ and dichloroethane. The sealed 210Pb-210Po sources were prepared directly by dropping an aliquot of the solution into the PPD cap followed by its evaporation. The sealed sources thus prepared were subjected to 1 m height drop test, air blowing test, and H₂O immersion

test. No radioactive contaminants were coming off from the sealed source through these tests.

CC 71-6 (Nuclear Technology)

ST alpha particle source polonium sealing
polycarbonate membrane

IT Membrane filters
(for preparation of low level sealed 210Pb source for random pulse generator)

IT Polycarbonates, uses
RL: NUU (Other use, unclassified); USES (Uses)
(for preparation of low level sealed 210Pb source for random pulse generator)

IT Radiation sources
Sealing
(preparation of low level sealed 210Pb source for random pulse generator)

IT 67-63-0, Isopropyl alcohol, uses 75-09-2, Dichloromethane, uses
1300-21-6, Dichloroethane 12027-17-7, Polonium hydroxide
(PO(OH)₄) 127795-35-1, Polonium nitrate
RL: NUU (Other use, unclassified); USES (Uses)
(for preparation of low level sealed 210Pb source for random pulse generator)

IT 12587-46-1, Alpha ray
RL: FMU (Formation, unclassified); FORM (Formation, nonpreparative)
(preparation of low level sealed & source
for random pulse generator)

IT 13981-52-7, Polonium-210, uses 14255-04-0, Lead-210,
uses
RL: TEM (Technical or engineered material use); USES (Uses)
(preparation of low level sealed 210Pb source for random pulse generator)

L84 ANSWER 8 OF 31 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2002:893088 CAPLUS Full-text
DOCUMENT NUMBER: 138:261764
TITLE: In vitro cell irradiation systems based on 210Po alpha
source: construction and characterization
AUTHOR(S): Szabo, J.; Feher, I.; Palfalvi, J.; Balashazy, I.;
Dam, A. M.; Polonyi, I.; Bogdandi, E. N.
CORPORATE SOURCE: KFKI Atomic Energy Research Institute, Budapest,
H-1525, Hung.
SOURCE: Radiation Measurements (2002), 35(6), 575-578
CODEN: RMEAEP; ISSN: 1350-4487
PUBLISHER: Elsevier Science Ltd.
DOCUMENT TYPE: Journal
LANGUAGE: English
AB One way of studying the risk to human health of low-level radiation exposure
is to make biol. expts. on living cell cultures. Two 210Po α -particle
emitting devices, with 0.5 and 100 MBq activity, were designed and constructed
to perform such expts. irradiating monolayers of cells. Ests. of dose rate at
the cell surface were obtained from measurements by a PIPS α -particle
spectrometer and from calcns. by the SRIM 2000, Monte Carlo charged particle
transport code. Particle fluence area distributions were measured by solid
state nuclear track detectors. The design and dosimetric characterization of
the devices are discussed.
CC 71-7 (Nuclear Technology)
Section cross-reference(s): 8
IT Animal tissue culture
Dosimeters

Radiation sources

(210Po α -particle emitting devices: design and dosimetric characterization for in vitro radiobiol. studies)

IT 13981-52-7, 210Po, uses

RL: BUU (Biological use, unclassified); DEV (Device component use); BIOL (Biological study); USES (Uses)

(210Po α -particle emitting devices: design and dosimetric characterization for in vitro radiobiol. studies)

IT 7440-69-9, Bismuth-209, uses 14255-04-0, 210Pb, uses

14331-79-4, 210Bi, uses

RL: PEP (Physical, engineering or chemical process); PYP (Physical process); TEM (Technical or engineered material use); PROC (Process); USES (Uses)

(210Po α -particle emitting devices: design and dosimetric characterization for in vitro radiobiol. studies)

REFERENCE COUNT: 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L84 ANSWER 9 OF 31 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 1997:743658 CAPLUS Full-text

DOCUMENT NUMBER: 128:67547

ORIGINAL REFERENCE NO.: 128:13107a,13110a

TITLE: A new method for evaluating annual alpha and beta dose rates in different ceramic samples by using solid state nuclear track detectors

AUTHOR(S): Misdaq, M. A.; Fahde, K.; Erramli, H.; Mikdad, A.

CORPORATE SOURCE: Nuclear Physics and Techniques Laboratory, Faculty of Sciences Semlalia, University Cadi Ayyad, Marrakech, Morocco

SOURCE: Radiation Physics and Chemistry (1997), 50(3), 293-297
CODEN: RPCHDM; ISSN: 0969-806X

PUBLISHER: Elsevier Science Ltd.

DOCUMENT TYPE: Journal

LANGUAGE: English

AB Uranium and thorium contents in different ceramic samples have been evaluated by using CR-39 and LR-115 type II solid state nuclear track detectors (SSNTDs) and calculating the probabilities for α -particles to reach and be registered on the SSNTD films. A new method has been developed based on calculating the stopping powers of the studied materials for the α - and β -particles emitted by the nuclei of the uranium and thorium series for evaluating annual α - and β -dose rates in the considered ceramic samples. β -Dose rates due to potassium 40 (40K) have been evaluated for the studied materials. α -Dose rates of the considered ceramic samples have been compared with data obtained by using Bell's method.

CC 71-7 (Nuclear Technology)

Section cross-reference(s): 57, 79

IT 25656-90-0, Diethylene glycol bis(allyl carbonate) homopolymer

RL: DEV (Device component use); USES (Uses)

(CR-39; evaluating annual α - and β -dose rates

emitted by U and Th series nuclei in different ceramic samples by using solid state nuclear track detectors)

IT 9004-70-0, LR 115II

RL: DEV (Device component use); USES (Uses)

(LR-115 type II; evaluating annual α - and β -dose

rates emitted by U and Th series nuclei in different ceramic samples by using solid state nuclear track detectors)

IT 7440-29-1, Thorium, analysis 14274-82-9, Thorium 228, analysis

14913-49-6, Bismuth 212, analysis 15389-34-1, Polonium 212, analysis

15756-58-8, Polonium 216, analysis 22481-48-7, Radon 220, analysis 28522-22-7, Radon 224, analysis
 RL: ANT (Analyte); ANST (Analytical study)
 (evaluating annual α -dose rates emitted by Th series nuclei in different ceramic samples by using solid state nuclear track detectors)

IT 7440-61-1, Uranium, analysis 13966-29-5, Uranium 234, analysis 13981-52-7, Polonium 210, analysis 13982-63-3, Radium 226, analysis 14269-63-7, Thorium 230, analysis 14859-67-7, Radon 222, analysis 15422-74-9, Polonium 218, analysis 15735-67-8, Polonium 214, analysis
 RL: ANT (Analyte); ANST (Analytical study)
 (evaluating annual α -dose rates emitted by U series nuclei in different ceramic samples by using solid state nuclear track detectors)

IT 14331-83-0, Actinium 228, analysis 14913-50-9, Thallium 208, analysis 15092-94-1, Lead 212, analysis 121239-98-3, Radon 228, analysis
 RL: ANT (Analyte); ANST (Analytical study)
 (evaluating annual β -dose rates emitted by Th series nuclei in different ceramic samples by using solid state nuclear track detectors)

IT 14255-04-0, Lead 210, analysis 14331-79-4, Bismuth 210, analysis 14733-03-0, Bismuth 214, analysis 15035-09-3, Thallium 206, analysis 15065-10-8, Thorium 234, analysis 15067-28-4, Lead 214, analysis 15100-28-4, Protactinium 234, analysis 51634-37-8, Thallium 218, analysis
 RL: ANT (Analyte); ANST (Analytical study)
 (evaluating annual β -dose rates emitted by U series nuclei in different ceramic samples by using solid state nuclear track detectors)

REFERENCE COUNT: 14 THERE ARE 14 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L84 ANSWER 10 OF 31 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 1983:412475 CAPLUS Full-text
 DOCUMENT NUMBER: 99:12475
 ORIGINAL REFERENCE NO.: 99:1947a,1950a
 TITLE: Investigation of nitric acid for removal of noxious radionuclides from uranium ore or mill tailings
 AUTHOR(S): Ryon, A. D.; Bond, W. D.; Hurst, F. J.; Scheitlin, F. M.; Seeley, F. G.
 CORPORATE SOURCE: Oak Ridge Natl. Lab., Oak Ridge, TN, 37830, USA
 SOURCE: Uranium Mill Tailings Manage., Proc. Two NEA Workshops (1982), Meeting Date 1981, 139-47. OECD: Paris, Fr.
 CODEN: 49VJAB
 DOCUMENT TYPE: Conference
 LANGUAGE: English
 AB A conceptual process using HNO₃ [7697-37-2] to extract 226Ra and 230Th in addition to the U from ore is proposed to decrease the potential hazard from discharge of mill tailings to the environment. This process removes $\leq 98\%$ of the 226Ra and 230Th, yielding a residue containing as low as 10 pCi of Ra/g. Leaching of U from ores is consistently $> 99.5\%$. The residue after multistage leaching with HNO₃ is resistant to further Ra leaching with water. Rn emanation from HNO₃-leached residues generally is low due to the low Ra content. Heating to 800° causes further reduction of Rn emanation. Greater than 99% recovery of Ra from HNO₃-leach solns. is obtained by carrying it on BaSO₄. Good adsorption of Ra is also obtained on barite and Celite. Recovery of Th and U by solvent extraction using Bu₃PO₄ appears promising. Recycle of HNO₃ may be accomplished by solvent extraction combined with evaporation and calcination.
 CC 71-10 (Nuclear Technology)

Section cross-reference(s): 54
 IT 10043-92-2, properties
 RL: PRP (Properties)
 (emanation of, from nitric acid leach solution of uranium ore)
 IT 7440-61-1P, properties 13981-52-7P, properties 13982-63-3P,
 properties 14255-04-0P, properties 14269-63-7P, properties
 RL: RCT (Reactant); PREP (Preparation); RACT (Reactant or reagent)
 (leaching of, from uranium ore, in nitric acid solution)

L84 ANSWER 11 OF 31 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 1978:185371 CAPLUS Full-text
 DOCUMENT NUMBER: 88:185371
 ORIGINAL REFERENCE NO.: 88:29107a,29110a
 TITLE: Analysis of alpha emitters in the
 coral, Favites virens, from Bikini Lagoon by
 solid-state track detection
 AUTHOR(S): Levy, Yitzhak; Miller, Donald S.; Friedman, Gerald M.;
 Noshkin, Victor E.
 CORPORATE SOURCE: Dep. Geol., Rensselaer Polytech. Inst., Troy, NY, USA
 SOURCE: Health Physics (1978), 34(3), 209-17
 CODEN: HLTPAO; ISSN: 0017-9078
 DOCUMENT TYPE: Journal
 LANGUAGE: English

AB A quant. method for the nondestructive anal. of α -particle emitters in CaCO_3 matrixes by solid-state track detection (cellulose nitrate) was developed. In an area of 4 mm^2 , .apprx.0.4 pCi/g can be measured routinely; smaller concns. can be determined but with a lower resolution. Calibration methods used were (a) a Pu source of 0.15 μCi in conjunction with polycarbonate and CaCO_3 absorbers of different thickness (2-30 μm) and (b) a powdered coral sample from Enewetak Atoll that had been radiochem. analyzed for Pu radionuclides, ^{241}Am , and other long-lived fission and activation products. Slabs of a coral, *F. virens*, from Bikini lagoon were analyzed. A quantity of the α -particle emitters detected in regions of the coral identified with growth during nuclear testing (1954, 1956, and 1959) are found in small discrete spots. Thin sections cut parallel to the direction of coral growth give different patterns of distributions. No such hot spots are evident in any post-test-yr growth sections although Pu and other long-lived fission and activation products were measured in these sections by radiochem. techniques.

CC 8-1 (Radiation Biochemistry)
 ST alpha particle emitter detn coral
 IT Favites virens
 (alpha-particle emitters determination in, by solid-state
 track detector)
 IT 7440-29-1, analysis 7440-61-1, analysis 13981-16-3, analysis
 13981-52-7, analysis 13982-63-3, analysis 14119-33-6, analysis
 14255-04-0, analysis 14269-63-7, analysis 14274-82-9, analysis
 14596-10-2, analysis 15117-48-3, analysis 15117-96-1, analysis
 RL: ANT (Analyte); ANST (Analytical study)
 (determination of, by solid-state track detector)

L84 ANSWER 12 OF 31 CAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 1977:570658 CAPLUS Full-text
 DOCUMENT NUMBER: 87:170658
 ORIGINAL REFERENCE NO.: 87:26955a,26958a
 TITLE: Emission and enrichments of radon daughters from Etna
 volcano magma
 AUTHOR(S): Lambert, G.; Bristeau, P.; Polian, G.
 CORPORATE SOURCE: Cent. Faibles Radioact., CNRS-CEA, Fr.
 SOURCE: Geophysical Research Letters (1976), 3(12), 724-6

CODEN: GPRLAJ; ISSN: 0094-8276

DOCUMENT TYPE: Journal
 LANGUAGE: English

AB Studies of the Etna volcano plume show that Rn daughters as well as gaseous Rn are directly emitted from the magma. Enrichment factors observed are 4 for Bi vs. Pb and 6 for Po vs. Pb. These enrichments correlate with relative volatility of the elements and/or their different compds. The ^{210}Pb total output measured for the Etna plume is an insignificant source for this nuclide.

CC 53-3 (Mineralogical and Geological Chemistry)
 IT 10043-92-2P, preparation 13981-52-7P, preparation
 14255-04-0P, preparation
 RL: PREP (Preparation)
 (emission of, from Etna Volcano)

L84 ANSWER 13 OF 31 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 1958:14581 CAPLUS Full-text
 DOCUMENT NUMBER: 52:14581
 ORIGINAL REFERENCE NO.: 52:2594a-b
 TITLE: Energy of α -particles from polonium-210
 AUTHOR(S): Agapkin, I. I.; Gol'din, L. L.
 SOURCE: Izvestiya Akademii Nauk SSSR, Seriya Fizicheskaya
 (1957), 21, 909-12
 CODEN: IANFAY; ISSN: 0367-6765
 DOCUMENT TYPE: Journal
 LANGUAGE: Unavailable

AB The measurements were made with a magnetic α -spectrometer. Error sources were listed; it was observed that freshly made prepns. had reproducible results. By using a correction for the half-width of the line, obtained by varying the slit width, an energy of 5297.8 ± 1.5 e.kv. was obtained, compared to the tabulated value of 5300.6 ± 2.6 e.kv. For $\text{Em}220$ the energy was 6282.4 (tabulated value 6282.3 ± 1.3 e.kv.).

CC 3A (Nuclear Phenomena)
 IT 13981-52-7P, Polonium, isotope of mass 210
 RL: PREP (Preparation)
 (separation from $\text{Bi}210$ and $\text{Pb}210$)
 IT 14255-04-0P, Lead, isotope of mass 210
 RL: PREP (Preparation)
 (separation from $\text{Bi}210$ and $\text{Po}210$)
 IT 13981-52-7, Polonium, isotope of mass 210 22481-48-7, Radon, isotope of mass 220
 (α -rays from)

L84 ANSWER 14 OF 31 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 1956:3794 CAPLUS Full-text
 DOCUMENT NUMBER: 50:3794
 ORIGINAL REFERENCE NO.: 50:690f-g
 TITLE: Purification of radioactive deposits formed in radon needles
 AUTHOR(S): Ferreira de Miranda, C.
 CORPORATE SOURCE: Fac. cienc., Lisbon, Port.
 SOURCE: Rev. fac. cienc. Univ. Lisboa (1954), 3, 191-5
 DOCUMENT TYPE: Journal
 LANGUAGE: Unavailable

AB Rn needles are treated with 3N HNO_3 and the liquid is centrifuged. The solution is heated up to 45° and a solution of 1.5N KCN is added. The precipitate is separated by centrifugation and redissolved in 0.5N HNO_3 . The procedure is repeated twice. The radioactive material formed by Ra D, Ra E, and Ra F is concentrated in the last HNO_3 solution

CC 3A (Nuclear Phenomena)
 IT 13981-52-7P, Polonium, isotope of mass 210 14255-04-0P,
 Lead, isotope of mass 210
 RL: PREP (Preparation)
 (concentration from Rn deposits)
 IT 10043-92-2, Radon
 (decay products of)

L84 ANSWER 15 OF 31 WPIX COPYRIGHT 2008 THOMSON REUTERS on STN
 DUPLICATE 6
 ACCESSION NUMBER: 1981-42085D [23] WPIX
 TITLE: Locating deposits of uranium and thorium - by collecting field samples forming chemical sons. and examining alpha sensitive films exposed to delay prods.
 DERWENT CLASS: K08; S03
 INVENTOR: STIEFF L R
 PATENT ASSIGNEE: (STIEFF-I) STIEFF L R
 COUNTRY COUNT: 2

PATENT INFORMATION:

| PATENT NO | KIND | DATE | WEEK | LA | PG | MAIN IPC |
|------------|------|----------|-----------|----|----|----------|
| US 4268748 | A | 19810519 | (198123)* | EN | | |
| CA 1145067 | A | 19830419 | (198319) | EN | | |

APPLICATION DETAILS:

| PATENT NO | KIND | APPLICATION | DATE |
|--------------|------|----------------|----------|
| US 4268748 A | | US 1978-879578 | 19780221 |
| US 4268748 A | | US 1978-899937 | 19780425 |
| US 4268748 A | | US 1979-106424 | 19791221 |

INT. PATENT CLASSIF.:

IPC RECLASSIF.: G01V0005-00 [I,A]; G01V0005-00 [I,C]
 ECLA: G01V0005-00
 USCLASS NCLM: 250/255.000
 NCLS: 250/472.100; 250/DIG.002

BASIC ABSTRACT:

US 4268748 A UPAB: 20050419 Uranium deposits are detected by collecting subsurface samples from a location and examining the samples for characteristic daughter prods. Pb, Bi and Po are chemically extracted from the samples. First and second alpha particle sensitive films are exposed to record decay of Po 214 and Po210. After etching, alpha particle damage populations are determined and correlated with sample locations in order to determine the position of a uranium deposit. Samples are collected from differentiated zones surrounding the deposit. Planchets on which alpha active Po from solution has been deposited are exposed to the alpha sensitive film. The method detects deposits of uranium by measuring Pb214, Bi 214, Po214, Pb210, and Po210 daughters. The deposits are distinguished from thorium deposits which can also be located. MANUAL CODE: CPI: K08-A; K09-J

EPI: S03-C03; S03-G01

L84 ANSWER 16 OF 31 WPIX COPYRIGHT 2008 THOMSON REUTERS on STN
 ACCESSION NUMBER: 2003-776157 [73] WPIX
 CROSS REFERENCE: 2003-197783; 2003-898376

DOC. NO. CPI: C2003-213524 [73]
 DOC. NO. NON-CPI: N2003-621799 [73]
 TITLE: Removal of radioactive contaminants from surface involves retaining aqueous solution comprising wetting agent and active agent on surface and removing aqueous solution containing radioactive contaminants from surface
 DERWENT CLASS: A17; A25; A97; K07; P43
 INVENTOR: MARTIN R T
 PATENT ASSIGNEE: (BOBO-N) BOBOLINK INC
 COUNTRY COUNT: 1

PATENT INFORMATION:

| PATENT NO | KIND | DATE | WEEK | LA | PG | MAIN IPC |
|------------|------|----------|-----------|----|------|----------|
| US 6605158 | B1 | 20030812 | (200373)* | EN | 8[0] | |

APPLICATION DETAILS:

| PATENT NO | KIND | APPLICATION | DATE |
|------------|-----------|----------------|----------|
| US 6605158 | B1 CIP of | US 2001-976467 | 20011012 |
| US 6605158 | B1 | US 2002-283039 | 20021024 |

FILING DETAILS:

| PATENT NO | KIND | PATENT NO |
|------------|-----------|--------------|
| US 6605158 | B1 CIP of | US 6497769 B |

PRIORITY APPLN. INFO: US 2002-283039 20021024
 US 2001-976467 20011012

INT. PATENT CLASSIF.:
 IPC RECLASSIF.: C11D0011-00 [I,A]; C11D0011-00 [I,C]; C11D0007-02 [I,C];
 C11D0007-08 [I,A]; C11D0007-22 [I,C]; C11D0007-26 [I,A];
 C11D0007-32 [I,A]; G21F0009-28 [I,A]; G21F0009-28 [I,C]
 ECLA: C11D0007-08; C11D0007-26E; C11D0007-32D; C11D0011-00B2D;
 C11D0011-00B8; G21F0009-28

ICO: M11D0007:26A
 USCLASS NCLM: 134/010.000
 NCLS: 134/002.000; 134/003.000; 134/022.190; 134/026.000;
 134/028.000; 134/036.000; 134/041.000; 134/042.000;
 210/682.000; 210/688.000; 376/308.000; 376/309.000;
 376/310.000; 423/002.000; 423/003.000; 423/020.000;
 510/110.000; 588/001.000; 588/020.000; 976/DIG.376;
 976/DIG.391

BASIC ABSTRACT:

US 6605158 B1 UPAB: 20050601

NOVELTY - Radioactive contaminants are removed from a surface by applying to the surface an aqueous solution comprising a wetting agent and an active agent; retaining the aqueous solution on the surface for a time to allow radioactive contaminants to migrate into the solution; and removing the solution containing the radioactive contaminants from the surface. The active agent is a specified complex substituted keto-amine.

DETAILED DESCRIPTION - Removal of radioactive contaminants from a surface includes applying to the surface an aqueous solution comprising a wetting agent and an active agent; retaining the aqueous solution on the surface for a time to allow radioactive contaminants to migrate into the aqueous solution; and removing the aqueous solution containing the radioactive contaminants from the surface. The wetting agent is methanol, ethanol, propanol, isopropanol,

butanol, propargyl alcohol, tertiary butyl alcohol, pentanol, propylene glycol, polypropylene glycol, and/or ethylene glycol. The active agent is a complex substituted keto-amine having the formula (I).



R = abietyl, hydroabietyl, or dehydroabietyl; R'' = alpha ketonyl having fewer than 10C; and R' = H, or CH_2R'' .

USE - For removing radioactive contaminants from a surface.

ADVANTAGE - The inventive method has the effect of removing all of the radioactive contaminants from a previously contaminated surface. It is simple, economic, and an effective way of removing all of the radioactive contaminants from various surfaces or areas. MANUAL CODE: CPI: A12-W11C; A12-W11F; K07-A03

ABEX EXAMPLE - Several decontamination solutions were prepared and tested utilizing various acids but containing the amount of keto-amine (0.5%), isopropyl alcohol (0.3%) and propargyl alcohol (0.3%). Each solution had a pH of at most 1.2. The solutions were applied as a low-pressure spray, allowed to remain at the site for 5 minutes and then removed by a squeegee. The spent solution was neutralized to a pH of 7, using a 50% sodium hydroxide solution. Surface contamination levels of both alpha and beta emissions were measured before and after treatment. The results showed degrees of effectiveness in removing radioactivity from surfaces utilizing a strong acid combined with a keto-amine as the active agent and a mixture of lower alcohols.

TECH

INORGANIC CHEMISTRY - Preferred Component: The aqueous solution further comprises an acid from hydrochloric acid, hydrofluoric acid, sulfuric acid, phosphoric acid, sulfurous acid, bromic acid, iodic acid, nitric acid, perchloric acid, oxalic acid, aqua regia, citric acid, sulfamic acid, glycolic acid, and/or ascorbic acid. It contains 0.1-1 wt.% wetting agent, 0.1-2 wt.% complex substituted keto-amine, and 0 wt.% acid. The complex substituted keto-amine has the molecular formula $C_{33}H_{45}NO_2C_1H$. The alpha ketonyl group is a ketone from acetone, methyl ethyl ketone, diacetone alcohol, isophorone, mesityl oxide, pentane dione, acetyl acetone, cyclopentanone, cyclohexanone, or acetophenone. The wetting agent is a mixture of isopropanol and propargyl alcohol.

INORGANIC CHEMISTRY - Preferred Component: The radioactive contaminant is a member of the lanthanide or actinide group. It is preferably Actinium-227, Americium-241, Americium-243, Antimony-124, Antimony-125, Barium-133, Beryllium-7, Bismuth-207, Cadmium-109, Calcium-45, Carbon-14, Cerium-139, Cerium-141, Cerium-144, Cesium-134, Cesium-135, Cesium-137, Chromium-51, Cobalt-56, Cobalt-57, Cobalt-58, Cobalt-60, Copper-67, Curium-242, Curium-243, Curium-244, Curium-247, Europium-152, Europium-154, Europium-155, Gadolinium-153, Germanium-68, Gold-195, Hafnium-181, Hydrogen-3 (Tritium), Iodine-125, Iodine-126, Iodine-129, Iodine-131, Iodine-133, Iridium-192, Iron-55, Iron-59, Lead-210, Manganese-54, Mercury-203, Neptunium-237, Nickel-59, Nickel-63, Niobium-94, Plutonium-236, Plutonium-238, Plutonium-239, Plutonium-240, Plutonium-241, Plutonium-242, Plutonium-243, Plutonium-244, Polonium-210, Potassium-40, Promethium-147, Protactinium-231, Radium-223, Radium-224, Radium-226, Radium-228, Ruthenium-106, Samarium-151, Scandium-46, Selenium-75, Silver-108m, Silver-110m, Sodium-22, Strontium-85, Strontium-89, Strontium-90, Sulfur-35, Tantalum-182, Technetium-99, Thallium-204, Thorium-natural, Thorium-228, Thorium-230, Thorium-232, Tin-113, Uranium-232, Uranium-233, Uranium-234, Uranium-235, Uranium-236, Uranium-238, Uranium-natural, Uranium-depleted, Yttrium-88, Yttrium-91, Zinc-65, Zirconium-95, or their associated decay products.

INSTRUMENTATION AND TESTING - Preferred Method: The applying, retaining and removing steps are repeated to optimize decontamination. The aqueous solution is applied to the surface by spraying. It is retained on the surface for less than 10 minutes.

Preferred Component: The surface is a metal, plastic, glass, concrete, wood, fiberglass, fabric, and/or soil.

L84 ANSWER 17 OF 31 BIOSIS COPYRIGHT (c) 2008 The Thomson Corporation on
STN DUPLICATE 4

ACCESSION NUMBER: 1997:358484 BIOSIS Full-text
DOCUMENT NUMBER: PREV199799664887

TITLE: Retrospective assessment of radon exposure by measurements of ^{210}Po implanted in surfaces using an alpha track detector technique.

AUTHOR(S): Falk, R.; Mellander, H.; Nyblom, L.; Ostergren, I.

CORPORATE SOURCE: Swedish Radiation Protection Inst., S-171 16 Stockholm, Sweden

SOURCE: Environment International, (1996) Vol. 22, No. SUPPL. 1, pp. S857-S861.
CODEN: ENVIDV. ISSN: 0160-4120.

DOCUMENT TYPE: Article
LANGUAGE: English
ENTRY DATE: Entered STN: 25 Aug 1997
Last Updated on STN: 25 Aug 1997

ABSTRACT: The radon exposure of the past is important in epidemiological studies where an assessment of lung cancer risk from indoor ***radon*** exposure is evaluated. The long-lived decay product, *** ^{210}Pb *** ($T/2 = 22$ y), is implanted into indoor surfaces by alpha recoils and can be monitored to give information about the previous radon history. This gives an alternative or complementary method to the traditional measurements of the current average radon concentration.

Autoradiographic alpha-track methods to assess the ^{210}Pb activity implanted in glass surfaces by measurement of ^{210}Po alpha activity were investigated to find a simple and reliable method for field use. One limiting factor at low exposure levels is the alpha background activity in the sub-surface material. In the search for a practical field method, the use of two different alpha-track detector materials was found successful. By exposing one Kodak LR-115 cellulose-nitrate film and one CR-39 detector side by side on glass panes, the background of the glass is measured with the LR-115 and both the background and the signal by the CR-39 detector. Results from measurements in 31 dwellings show that an exposure of more than 1000 Bq m^{-3} to a glass surface can be measured with the (CR-LR) difference technique. Experiences from the field measurements show the method to be accurate, simple, and reliable and therefore a promising tool for future radon epidemiological studies.

CONCEPT CODE: Radiation biology - Radiation effects and protective measures 06506
Biochemistry studies - Minerals 10069
Toxicology - Environment and industry 22506
Public health - Air, water and soil pollution 37015
Public health - Radiation health 37017

INDEX TERMS: Major Concepts
Biochemistry and Molecular Biophysics; Pollution Assessment Control and Management; Radiation Biology; Toxicology

INDEX TERMS: Chemicals & Biochemicals
RADON; POLONIUM-210; CR-39

INDEX TERMS: Miscellaneous Descriptors
ALPHA TRACK DETECTOR TECHNIQUE; ASSESSMENT METHOD; CARCINOGEN; CR-39 DETECTOR; EPIDEMIOLOGICAL STUDIES; EXPOSURE; INDOOR; LONG-LIVED DECAY PRODUCT; LUNG CANCER;

NEOPLASTIC DISEASE; POLLUTION; POLONIUM-210; RADIATION BIOLOGY; RADIONUCLIDES; RADON; RESPIRATORY SYSTEM DISEASE; RISK

REGISTRY NUMBER: 10043-92-2 (RADON)
 13981-52-7 (POLONIUM-210)
 25656-90-0Q (CR-39)
 81283-55-8Q (CR-39)

L84 ANSWER 18 OF 31 BIOSIS COPYRIGHT (c) 2008 The Thomson Corporation on
 STN DUPLICATE 5

ACCESSION NUMBER: 1989:159461 BIOSIS Full-text
 DOCUMENT NUMBER: PREV198987081562; BA87:81562
 TITLE: POLONIUM IN FLORIDA USA GROUNDWATER AND ITS POSSIBLE
 RELATIONSHIP TO THE SULFUR CYCLE AND BACTERIA.
 AUTHOR(S): HARADA K [Reprint author]; BURNETT W C; LAROCK P A; COWART
 J B
 CORPORATE SOURCE: DEP OCEANOGRAPHY, FLA STATE UNIV, TALLAHASSEE, FLA 32306,
 USA
 SOURCE: Geochimica et Cosmochimica Acta, (1989) Vol. 53, No. 1, pp.
 143-150.
 CODEN: GCACAK. ISSN: 0016-7037.

DOCUMENT TYPE: Article
 FILE SEGMENT: BA
 LANGUAGE: ENGLISH
 ENTRY DATE: Entered STN: 25 Mar 1989
 Last Updated on STN: 25 Mar 1989

ABSTRACT: The last radioactive member of the 238U natural decay-series, ***210Po*** is normally considered a very particle-reactive isotope. Analysis of most natural waters shows that 210Po is present at very low activities, usually even lower than its insoluble precursor, 210Pb. We have recently discovered, however, that 210Pb exists at very high concentrations in groundwaters of some shallow aquifers in west central Florida. These waters tend to be fairly acidic (pH < 5), sulfide-bearing, and relatively high in 222Rn. Detailed study of one well with extraordinary levels of 210Pb (.apprx. 1000 dpm/l) indicates that: (1) 210Po in this water is in great excess of radioactive equilibrium with its predecessors ***210Pb*** and 210Bi; (2) most Po in this water exists in a form which does not coprecipitate with an iron hydroxide scavenger; and (3) the conversion of soluble (0.2 µm filter) to particulate Po occurs over a time scale of a few days during sulfide oxidation. We suspect that Po cycling in this environment is related to the sulfur cycle and may, therefore, be influenced by sulfur bacteria.

CONCEPT CODE: General biology - Conservation and resource management 00512
 Radiation biology - General 06502
 Circadian rhythms and other periodic cycles 07200
 Ecology: environmental biology - Oceanography and limnology 07510
 Biochemistry studies - Minerals 10069
 Bacteriology, general and systematic 30000
 Physiology and biochemistry of bacteria 31000
 Food microbiology - Biodegradation and biodeterioration 39006
 Soil science - Physics and chemistry 52805
 INDEX TERMS: Major Concepts
 Biochemistry and Molecular Biophysics;
 Biosynchronization; Conservation; Estuarine Ecology
 (Ecology, Environmental Sciences); Radiation Biology;
 Soil Science; Systematics and Taxonomy

INDEX TERMS: Miscellaneous Descriptors

ORGANISM: AQUIFER CHEMICAL ANALYSIS RADON URANIUM
 Classifier
 Bacteria 05000
 Super Taxa
 Microorganisms
 Taxa Notes
 Bacteria, Eubacteria, Microorganisms

REGISTRY NUMBER: 7440-08-6 (POLONIUM)
 7704-34-9 (SULFUR)
 10043-92-2 (RADON)
 7440-61-1 (URANIUM)

L84 ANSWER 19 OF 31 BIOSIS COPYRIGHT (c) 2008 The Thomson Corporation on
 STN

ACCESSION NUMBER: 2003:4191 BIOSIS Full-text
 DOCUMENT NUMBER: PREV200300004191

TITLE: Experimental methods of determining the activity depth distribution of implanted ^{210}Po in glass.

AUTHOR(S): Roos, Birgitta [Reprint Author]; Samuelsson, Christer

CORPORATE SOURCE: Department of Radiation Physics, Lund University Hospital,
 SE-221 85, Lund, Sweden
 birgitta.roos@radfys.lu.se

SOURCE: Journal of Environmental Radioactivity, (2002) Vol. 63, No.
 2, pp. 135-151. print.
 ISSN: 0265-931X (ISSN print).

DOCUMENT TYPE: Article
 LANGUAGE: English
 ENTRY DATE: Entered STN: 18 Dec 2002
 Last Updated on STN: 18 Dec 2002

ABSTRACT: Glass is often used in radon surveys to estimate retrospective radon concentrations, as radon progenies are embedded in the upper surface layer. Experimental methods based on etching to determine the depth distribution of recoil-implanted ^{210}Po in glass from radon decay in air is presented. By carefully controlling chemical concentrations and exposure time during which the glass is etched, stepwise removal of the surface material was possible. Two different etching agents, diluted HF/HNO₃ and NaOH were utilised, with very similar results. Experimental recoil depths of ^{210}Po agree with theoretical calculations from the literature. The maximum implantation depth obtained using this procedure was 100+-20nm.

CONCEPT CODE: Radiation biology - General 06502
 Biochemistry studies - General 10060

INDEX TERMS: Major Concepts
 Biochemistry and Molecular Biophysics; Radiation Biology
 INDEX TERMS: Chemicals & Biochemicals
 glass; hydrofluoric acid; lead-210;
 depth distribution; nitric acid; radon; sodium hydroxide

INDEX TERMS: Methods & Equipment
 radon survey: applied and field techniques

REGISTRY NUMBER: 7664-39-3 (hydrofluoric acid)
 14255-04-0 (lead-210)
 7697-37-2 (nitric acid)
 10043-92-2 (radon)
 1310-73-2 (sodium hydroxide)

L84 ANSWER 20 OF 31 BIOSIS COPYRIGHT (c) 2008 The Thomson Corporation on
 STN

ACCESSION NUMBER: 1985:424029 BIOSIS Full-text
 DOCUMENT NUMBER: PREV198580094021; BA80:94021

TITLE: A PRACTICAL METHOD FOR THE SIMULTANEOUS DETERMINATION OF
 THORIUM-234 RADON-226 LEAD-210
 AND POLONIUM-210 IN SEAWATER.
 AUTHOR(S): HARADA K [Reprint author]; TSUNOGAI S
 CORPORATE SOURCE: DEP CHEM, FACULTY FISHERIES, HOKKAIDO UNIV, HAKODATE, 041,
 JAPAN
 SOURCE: Journal of the Oceanographical Society of Japan, (1985)
 Vol. 41, No. 2, pp. 98-104.
 CODEN: NKGKB4. ISSN: 0029-8131.
 DOCUMENT TYPE: Article
 FILE SEGMENT: BA
 LANGUAGE: ENGLISH
 ABSTRACT: A practical method was developed for the simultaneous determination of 226Ra, 234Th, 210Pb and 210Po in seawater. Samples are spiked with 228Ra, 230Th, 208Po and common Pb to determine chemical yield. These nuclides are coprecipitated with calcium carbonate and ferric ***hydroxide*** from 20-50 l of seawater and separated by using coprecipitation and ion exchange techniques. Counting sources of Ra and the other nuclides are prepared by electrodeposition onto silver discs. Their radioactivities are counted with an α -spectrometer and a low background β -counter. This method gives a standard deviation of .apprx. 5% for replicate determination of 226Ra and other nuclides.
 CONCEPT CODE: Methods - Field methods 01008
 Mathematical biology and statistical methods 04500
 Radiation biology - Radiation and isotope techniques 06504
 Ecology: environmental biology - Oceanography 07512
 Biochemistry methods - General 10050
 Biochemistry studies - General 10060
 Biophysics - Methods and techniques 10504
 INDEX TERMS: Major Concepts
 Biochemistry and Molecular Biophysics; Marine Ecology
 (Ecology, Environmental Sciences); Methods and
 Techniques; Radiology (Medical Sciences)
 INDEX TERMS: Miscellaneous Descriptors
 COPRECIPITATION ION EXCHANGE ELECTRODEPOSITION
 RADIOACTIVITY SPECTROMETER
 REGISTRY NUMBER: 15065-10-8 (THORIUM-234)
 16369-95-2 (RADON-226)
 14255-04-0 (LEAD-210)
 13981-52-7 (POLONIUM-210)

L84 ANSWER 21 OF 31 ENERGY COPYRIGHT 2008 USDOE/IEA-ETDE on STN
 ACCESSION NUMBER: 2008(2):11838 ENERGY Full-text
 TITLE: Uncertainties of retrospective radon concentration
 measurements by multilayer surface trap detector.
 AUTHOR: Bastrikov, V.; Kruzhakov, A. (Ural State Technical
 Univ., Yekaterinburg (Russian Federation)); Zhukovsky,
 M. (Institute of Industrial Ecology UB RAS,
 Yekaterinburg (Russian Federation))
 CORPORATE SOURCE: Societe Francaise de Radioprotection - SFRP, BP72,
 92263 Fontenay-aux-Roses Cedex (France)
 NUMBER OF REPORT: INIS-FR--7038
 SOURCE: 2006. 11 p. Available from INIS in electronic form.
 Conference: Second European IRPA congress on radiation
 protection - Radiation protection: from knowledge to
 action, Paris (France), 15-19 May 2006
 DOCUMENT TYPE: Miscellaneous; Availability Note; Conference
 COUNTRY: France
 LANGUAGE: English

FIELD AVAILABILITY: AB

ABSTRACT: The detector for retrospective radon exposure measurements is developed. The detector consists of the multilayer package of solid-state nuclear track detectors LR-115 type. Nitrocellulose films works both as A-particle detector and as absorber decreasing the energy of A-particles. The uncertainties of implanted ^{210}Pb measurements by two- and three-layer detectors are assessed in dependence on surface ^{210}Po activity and gross background activity of the glass. The generalized compartment behavior model of radon decay products in the room atmosphere was developed and verified. It is shown that the most influencing parameters on the value of conversion coefficient from ^{210}Po surface activity to average radon concentration are aerosol particles concentration, deposition velocity of unattached ^{218}Po and air exchange rate. It is demonstrated that with the use of additional information on surface to volume room ratio, air exchange rate and aerosol particles concentration the systematic bias of conversion coefficient between surface activity of ^{210}Po and average radon concentration can be decreased up to 30 %. (N.C.)

CLASSIFICATION CODE: *S46 INSTRUMENTATION RELATED TO NUCLEAR SCIENCE AND TECHNOLOGY

CONTROLLED TERM: ALPHA DECAY RADIOISOTOPES; LEAD 210; LEAD 214; POLONIUM 214; POLONIUM 218; RADIATION DETECTORS; RADIATION DOSES; RADON; RISK ASSESSMENT; THORIUM 232; URANIUM 238

BROADER TERM: ACTINIDE NUCLEI; ALPHA DECAY RADIOISOTOPES; BETA DECAY RADIOISOTOPES; BETA-MINUS DECAY RADIOISOTOPES; DOSES; ELEMENTS; EVEN-EVEN NUCLEI; FLUIDS; GASES; HEAVY NUCLEI; ISOTOPES; LEAD ISOTOPES; MEASURING INSTRUMENTS; MICROSECONDS LIVING RADIOISOTOPES; MINUTES LIVING RADIOISOTOPES; NONMETALS; NUCLEI; POLONIUM ISOTOPES; RADIOISOTOPES; RARE GASES; SPONTANEOUS FISSION RADIOISOTOPES; THORIUM ISOTOPES; URANIUM ISOTOPES; YEARS LIVING RADIOISOTOPES

L84 ANSWER 22 OF 31 ENERGY COPYRIGHT 2008 USDOE/IEA-ETDE on STN

ACCESSION NUMBER: 2006(1):198 ENERGY Full-text

TITLE: ^{210}Po and ^{210}Pb in seals from the Baltic Sea and Lake Saimaa, Finland.

AUTHOR: Solatie, D.; Rissanen, K. (Radiation and Nuclear Safety Authority - STUK, Rovaniemi (Finland). Regional Laboratory in Northern Finland); Vesterbacka, P. (Radiation and Nuclear Safety Authority - STUK, Helsinki (Finland). Natural Radiation Laboratory) SSI--2005-15

NUMBER OF REPORT: SOURCE: Radiological Protection in Transition. Proceedings of the 14. Regular Meeting of the Nordic Society for Radiation Protection, NSFS.

Editor(s): Valentin, J.; Cederlund, T.; Drake, P.; Finne, I.E.; Glansholm, A.; Jaworska, A.; Paile, W.; Rahola, T.

Swedish Radiation Protection Authority, Stockholm (Sweden)

Sep 2005. p. 237 of 386 p. Available from:
http://www.ssi.se/ssi/sub_r/apporter/pdf/ssi/sub_r/app/sub_2/005/sub_1/5.pdf; OSTI; Commercial reproduction prohibited; OSTI as DE20674930; PURL: <https://www.osti.gov/servlets/purl/20674930-Up6xtr/>. Conference: Radiological Protection in Transition. 14. Regular Meeting of the Nordic Society for Radiation Protection, NSFS, Raettvik (Sweden), 27 - 31 Aug 2005

ISSN: 0282-4434

DOCUMENT TYPE: Report Article; Conference

COUNTRY: Sweden
 LANGUAGE: English
 FIELD AVAILABILITY: AB
 ABSTRACT: 210Po and 210Pb are members of the 238U decay chain. 210Po is an alpha-emitter with a half-life of 138 days, while its grandmother, 210Pb is a beta-emitter with 22.3 year half-life. In the atmosphere 222Rn formats its decay products 210Po and 210Pb. These nuclides are deposited on to the surface of land and sea and thus enter the food chain. The naturally occurring radionuclides 210Po and 210Pb are important because their great contribution to radiation dose to human and other species. As top predators in the aquatic food chain, fish-eating seals are vulnerable to the accumulation of contaminants. In the Regional Laboratory in Northern Finland, measurements of 210Po and 210Pb activity concentrations in seals from the Baltic Sea and in ringed seals from Lake Saimaa have been performed. Concentrations of 210Po and 210Pb in seals were determined in muscle, liver, kidney and spleen. The results of 210Po and 210Pb activity concentrations and the ratio of 210Po / 210Pb in these samples are presented. (Summary-only contribution)
 CLASSIFICATION CODE: *S54 Environmental sciences
 CONTROLLED TERM: PINNIPEDS; BALTIC SEA; LAKES; POLONIUM 210; LEAD 210; RADIOECOLOGICAL CONCENTRATION; MUSCLES; LIVER; SPLEEN; KIDNEYS
 BROADER TERM: ALPHA DECAY RADIOISOTOPES; ANIMALS; AQUATIC ORGANISMS; BETA DECAY RADIOISOTOPES; BETA-MINUS DECAY; RADIOISOTOPES; BODY; DAYS LIVING RADIOISOTOPES; DIGESTIVE SYSTEM; ECOLOGICAL CONCENTRATION; EVEN-EVEN NUCLEI; GLANDS; HEAVY NUCLEI; ISOMERIC TRANSITION ISOTOPES; ISOTOPES; LEAD ISOTOPES; MAMMALS; NANOSECONDS LIVING RADIOISOTOPES; NUCLEI; ORGANS; POLONIUM ISOTOPES; RADIOISOTOPES; SEAS; SURFACE WATERS; VERTEBRATES; YEARS LIVING RADIOISOTOPES
 ELEMENT TERM: Po; 210Po; is; Po is; Pb; 210Pb; Pb is; U; 238U; U is; Rn; 222Rn; Rn is

L84 ANSWER 23 OF 31 ENERGY COPYRIGHT 2008 USDOE/IEA-ETDE on STN
 ACCESSION NUMBER: 2004(14):81802 ENERGY Full-text
 TITLE: Preparation of low level sealed 210Pb source for random pulse generator.
 AUTHOR: Mitsugashira, T.; Hara, M. (Tohoku Univ., Institute for Materials Research, Oarai Branch, Oarai, Ibaraki (Japan)); Tsuyuzaki, N. (OPTRANS Corp. (Japan))
 NUMBER OF REPORT: KEK-PROC--2003-11
 SOURCE: Proceedings of the fourth workshop on environmental radioactivity.
 Editor(s): Miura, Taichi
 High Energy Accelerator Research Organization, Tsukuba, Ibaraki (Japan)
 Nov 2003. p. 263-268 of 384 p. 2 refs., 5 figs., 1 tab. Available from KEK(High Energy Accelerator Research Organization) 1-1 Oho, Tsukuba-shi, Ibaraki-ken, 305-0801 JAPAN.
 Conference: 4. workshop on environmental radioactivity, Tsukuba, Ibaraki (Japan), 4 - 6 Mar 2003
 DOCUMENT TYPE: Report Article; Conference; Availability Note
 COUNTRY: Japan
 LANGUAGE: Japanese
 FIELD AVAILABILITY: AB
 ABSTRACT: We have developed the random pulse generator (RPG) that utilizes alpha-particle detection with pin photodiode (PPD). In order to support an expected large market of RPG, a steady production system of weak (about 100 Bq) alpha sealed source is necessary, and, for such alpha-source, 210Pb-210Po source is the best

candidate on a viewpoint of environmental radioactivity impact. Two methods for such 210Pb-210Po sealed source preparation, namely direct deposition from isopropyl alcohol solution (IPA solution) of 210Pb-210Po nitrate (D-IPA method) and the 210Pb-210Po hydroxides precipitation (PPT method), were experimentally examined. In the former D-IPA method, an aliquot of IPA solution of 210Pb-210Po nitrate was directly dropped in a sealed cap for PPD and dried by heating. Then, a polycarbonate (PC) solution of 1/1 mixture of dichloromethane and dichloroethane was dropped on the source to make a thin (about 0.1 mg/cm²) film for radioactivity sealed. In the PPT method, 210Pb-210Po hydroxide was filtrated on a PC membrane filter (Nuclipore 0.1 μm) and the membrane filter was dissolved in a 1/1 mixture of dichloromethane and dichloroethane. The sealed 210Pb-210Po sources were prepared directly by dropping an aliquot of the solution into the PPD cap followed by its evaporation. The sealed sources thus prepared were subjected to 1m height fall-down test, air blowing test, and water immersion test. It was confirmed that no radioactive contaminants were coming off from the sealed source through these tests. (author)

CLASSIFICATION CODE: *S07 Isotopes and radiation sources
 CONTROLLED TERM: ALCOHOLS; ALPHA DETECTION; COMPARATIVE EVALUATIONS; DEPOSITION; FABRICATION; HYDROXIDES; LEAD 210; NITRATES; POLONIUM 210; PRECIPITATION; PULSE GENERATORS; SEALED SOURCES
 BROADER TERM: ALPHA DECAY RADIOISOTOPES; BETA DECAY RADIOISOTOPES; BETA-MINUS DECAY RADIOISOTOPES; CHARGED PARTICLE DETECTION; DAYS LIVING RADIOISOTOPES; DETECTION; ELECTRONIC EQUIPMENT; EQUIPMENT; EVALUATION; EVEN-EVEN NUCLEI; FUNCTION GENERATORS; HEAVY NUCLEI; HYDROGEN COMPOUNDS; HYDROXY COMPOUNDS; ISOMERIC TRANSITION ISOTOPES; ISOTOPES; LEAD ISOTOPES; NANOSEC LIVING RADIOISOTOPES; NITROGEN COMPOUNDS; NUCLEI; ORGANIC COMPOUNDS; OXYGEN COMPOUNDS; POLONIUM ISOTOPES; RADIATION DETECTION; RADIATION SOURCES; RADIOISOTOPES; SEPARATION PROCESSES; YEARS LIVING RADIOISOTOPES
 ELEMENT TERM: Pb; 210Pb; is; Pb is; Pb*Po; Pb sy 2; sy 2; Po sy 2; Po is; 210Po; 210Pb-210Po; D

L84 ANSWER 24 OF 31 ENERGY COPYRIGHT 2008 USDOE/IEA-ETDE on STN
 ACCESSION NUMBER: 1994(14):96992 ENERGY Full-text
 TITLE: Evaluation of radionuclide levels and radiological dose in three populations of marine mammals in the eastern Canadian Arctic.
 AUTHOR: Macdonald, C.R.; Ewing, L.L.; Wiewel, A.M.; Harris, D.A. (AECL Research, Pinawa, Manitoba (Canada)); Stewart, R.E.A. (Dept. of Fisheries and Oceans, Winnipeg, Manitoba (Canada). Freshwater Inst.)
 CONF-931152--
 NUMBER OF REPORT: Ecological risk assessment: Lessons learned?. Abstract book.
 SOURCE: Anon.
 Pensacola, FL: Society of Environmental Toxicology and Chemistry. 1993. p. 254 of 356 p. Society of Environmental Toxicology and Chemistry Office, 1010 North 12th Avenue, Pensacola, FL 32501-3307 (United States).
 Conference: 14. annual meeting of the Society of Environmental Toxicology and Chemistry (SETAC), Houston, TX (United States), 14-18 Nov 1993
 DOCUMENT TYPE: Book Article; Conference
 COUNTRY: United States
 LANGUAGE: English
 FIELD AVAILABILITY: AB

ABSTRACT: Radionuclide levels were measured in beluga, walrus and ringed seal populations collected in 1992 to assess radiation dose and changes in dose with age and sex. The authors hypothesized that Arctic marine food chains accumulate high levels of naturally-occurring radionuclides such as polonium-210 and that radiation may pose a stress to animals which also accumulate metals such as cadmium. Liver, kidney, muscle and jawbone were analyzed by gamma spectrometry for cesium-137, cesium-134, lead-210, potassium-40 and radium-226 and fission-derived nuclides. Polonium-210 was analyzed by alpha spec after autodeposition onto a silver disk. Cesium-137 concentrations in muscle in all three populations were low, and ranged from below detection limits to 10 Bq/kg ww. There was no evidence of fission-derived radionuclides such as zinc-65 or cobalt-60. Lead-210 levels ranged from below detection limits in muscle of ringed seal and walrus to a mean of 82.3 Bq/kg ww in walrus bone. Polonium-210 in the three population ranged from 10 to 30 Bq/kg ww in bone and kidney. The major contributor to dose in the animals was polonium-210 because it is an alpha emitter and accumulates to moderate levels in liver and kidney. Radiological dose is approximately 20--30 times higher than background in humans, and is considerably lower than the dose observed in terrestrial food chains in the Arctic

CLASSIFICATION CODE: *560162; C2120

CONTROLLED TERM: AQUATIC ORGANISMS; CESIUM 134; CESIUM 137; CONTAMINATION; FOOD CHAINS; LEAD 210; MAMMALS; NATURAL RADIOACTIVITY; POLONIUM 210; POTASSIUM 40; RADIUM 226; SENSITIVITY; TISSUE DISTRIBUTION
 *MAMMALS: *SENSITIVITY; *FOOD CHAINS: *CONTAMINATION;
 *CESIUM 137: *TISSUE DISTRIBUTION; *CESIUM 134:
 *TISSUE DISTRIBUTION; *LEAD 210: *TISSUE DISTRIBUTION;
 *POTASSIUM 40: *TISSUE DISTRIBUTION; *RADIUM 226:
 *TISSUE DISTRIBUTION; *POLONIUM 210: *TISSUE DISTRIBUTION

BROADER TERM: ALKA; ALKALI METAL ISOTOPES; ALKALINE EARTH ISOTOPES; ALPHA DECAY RADIOISOTOPES; ANIMALS; BETA DECAY RADIOISOTOPES; BETA-MINUS DECAY RADIOISOTOPES; BETA-PLUS DECAY RADIOISOTOPES; CARBON 14 DECAY RADIOISOTOPES; CESIUM ISOTOPES; DAYS LIVING RADIOISOTOPES; DISTRIBUTION; ELECTRON CAPTURE RADIOISOTOPES; EVEN-EVEN NUCLEI; HEAVY ION DECAY RADIOISOTOPES; HEAVY NUCLEI; HOURS LIVING RADIOISOTOPES; INTERMEDIATE MASS NUCLEI; INTERNAL CONVERSION RADIOISOTOPES; ISOMERIC TRANSI; ISOMERIC TRANSITION ISOTOPES; ISOTOPES; LEAD ISOTOPES; LIGHT NUCLEI; NANOSEC LIVING RADIOISOTOPES; NUCLEI; ODD-EVEN NUCLEI; ODD-ODD NUCLEI; POLONIUM ISOTOPES; POTASSIUM ISOTOPES; RADIOACTIVITY; RADIOISOTOPES; RADIUM ISOTOPES; VERTEBRATES; YEARS LIVING RADIOISOTOPES

L84 ANSWER 25 OF 31 ENERGY COPYRIGHT 2008 USDOE/IEA-ETDE on STN
 ACCESSION NUMBER: 1985(18):134853 ENERGY Ful-text
 TITLE: Application of dual NAI-CsI(Tl) detectors to the in vivo detection and localization of radon seeds.
 AUTHOR: Anon. [United States]
 CORPORATE SOURCE: New York Univ., NY (USA). Inst. of Environmental Medicine (4664000)
 NUMBER OF REPORT: DOE/EV/04326--6; DE85001413
 SOURCE: In vivo measurements of bone-seeking radionuclides. Progress report, 1981-1984.
 Cohen, N.
 1984. pp. 89-96 Availability: NTIS, PC A06/MF A01; 1.
 DOCUMENT TYPE: Report Article
 COUNTRY: United States
 LANGUAGE: English
 DOCUMENT NUMBER: ERA-10:043005

ABSTRACT: Sealed gold capillary tubes containing radon have been used as implants for the treatment of certain tumors and lesions. The short-lived daughters of Rn-222 decay with the 3.82 day half-life of the parent. The seeds are generally left after treatment since it is believed that the residual activity from Pb-210, Bi-210 and Po-210 does not represent a significant radiological hazard to the patient. Some have suggested that chronic low-level irradiation of the implantation site may result in the formation of a tumor. Radon seeds which were implanted more than 20 years ago are now being considered for removal. Three subjects were measured in our whole body counting facility using NaI-CsI(T1) detectors. With the NaI-CsI(T1) detector placed directly over the implantation area, it was possible to observe the K X-rays characteristic of gold which are produced as a result of beta particle interaction with the gold casing. 2 refs., 3 figs., 1 tab. CLASSIFICATION CODE: *560151; C1500

CONTROLLED TERM:

*RADIATION SOURCE IMPLANTS; *RADIATION HAZARDS; *RADON 222; *RADIATION SOURCE IMPLANTS; BISMUTH 210; BREMSSTRAHLUNG; DAUGHTER PRODUCTS; GOLD; JAW; LEAD 210; LEGS; MAMMARY GLANDS; NEOPLASMS; RADIOTHERAPY; SOLID SCINTILLATION DETECTORS; WHOLE-BODY COUNTERS; X RADIATION

BROADER TERM:

ALPHA DECAY RADIOISOTOPES; BETA DECAY RADIOISOTOPES; BETA-MINUS DECAY RADIOISOTOPES; BISMUTH ISOTOPES; BODY; BODY AREAS; DAYS LIVING RADIOISOTOPES; DISEASES; ELECTROMAGNETIC RADIATION; ELEMENTS; EVEN-EVEN NUCLEI; GLANDS; HAZARDS; HEALTH HAZARDS; HEAVY NUCLEI; IMPLANTS; IONIZING RADIATIONS; ISOTOPES; LEAD ISOTOPES; LIMBS; MEASURING INSTRUMENTS; MEDICINE; METALS; NUCLEAR MEDICINE; NUCLEI; ODD-ODD NUCLEI; ORGANS; RADIATION DETECTORS; RADIATION SOURCES; RADIATIONS; RADIOISOTOPES; RADIOLOGY; RADON ISOTOPES; SCINTILLATION COUNTERS; SKELETON; SKULL; THERAPY; TRANSITION ELEMENTS; YEARS LIVING RADIOISOTOPES

ELEMENT TERM:

Cs*I*Na*T; Cs sy 4; sy 4; I sy 4; Na sy 4; T sy 4; NaI; Na cp; cp; I cp; CsI(T; Cs cp; T cp; NaI-CsI(T; Rn; Pb; Bi; Po; K

L84 ANSWER 26 OF 31 ENERGY COPYRIGHT 2008 USDOE/IEA-ETDE on STN

ACCESSION NUMBER:

1983(10):83361 ENERGY Full-text

TITLE:

Field method for detecting deposits containing uranium and thorium. (Patent.)

AUTHOR:

Stieff, L.R. [United States]

PATENT INFORMATION:

US 4336451 22 Jun 1982

v p. PAT-APPL-106424.

APPLICATION INFORMATION: 21 Dec 1979

DOCUMENT TYPE: Patent

COUNTRY: United States

LANGUAGE: English

ABSTRACT:

Locations of buried deposits are determined by detecting the presence of ^{238}U , ^{234}Pa , ^{234}Bi , ^{234}Po , ^{210}Po , ^{210}Bi and ^{210}Po in solutions obtained by chemically leaching these elements from rocks and soil. Polonium from the solution is plated on silver foil planchets. Alpha sensitive films are exposed to the alpha decay of ^{234}Po and ^{210}Po by contacting the films with the planchets. The films, when etched, reveal the damage caused by the passage of the high energy alpha particles. Alpha damage as a function of sample size, volume of solution used, planchet, foil or film area and exposure times measures concentration of ^{234}Po and ^{210}Po in the sample. Anomalous concentrations suggest presence of buried deposits containing uranium. Similar anomalous

concentrations of alpha damage from 212Bi and 212Po formed in films exposed to foils or planchets plated from leached solutions containing 212Pb, 212Bi and 212Po suggest deposits containing thorium. Plotting normalized alpha damage population (Numbers of alpha per gram, per square mm., per hour) and sample locations suggest mineral deposit locations.

INT. PATENT CLASSIF.: G01V005-00
 CLASSIFICATION CODE: *050200; 050100; B3100
 CONTROLLED TERM: *URANIUM DEPOSITS; *GEOPHYSICAL SURVEYS; ALPHA DETECTION; BISMUTH 210; BISMUTH 214; LEACHATES; LEAD 210; LEAD 214; POLONIUM 210; POLONIUM 214; QUANTITY RATIO; THORIUM
 BROADER TERM: ACTINIDES; ALPHA DECAY RADIOISOTOPES; BETA DECAY RADIOISOTOPES; BETA-MINUS DECAY RADIOISOTOPES; BISMUTH ISOTOPES; CHARGED PARTICLE DETECTION; DAYS LIVING RADIOISOTOPES; DETECTION; DISPERSIONS; ELEMENTS; EVEN-EVEN NUCLEI; GEOLOGIC DEPOSITS; HEAVY NUCLEI; ISOTOPES; LEAD ISOTOPES; METALS; MICROSEC LIVING RADIOISOTOPES; MINERAL RESOURCES; MINUTES LIVING RADIOISOTOPES; MIXTURES; NUCLEI; ODD-ODD NUCLEI; POLONIUM ISOTOPES; RADIATION DETECTION; RADIOISOTOPES; RESOURCES; SECONDS LIVING RADIOISOTOPES; SOLUTIONS; SURVEYS; YEARS LIVING RADIOISOTOPES
 ELEMENT TERM: B*I; BI; 214BI; is; B is; 214B; B cp; cp; I cp; O*P; PO; 214PO; P is; 214P; P cp; O cp; B*P; BP; 210BP; 210B; 210BI; 210PO; 210P; 212BI; 212B; 212PO; 212P

L84 ANSWER 27 OF 31 ENERGY COPYRIGHT 2008 USDOE/IEA-ETDE on STN
 ACCESSION NUMBER: 1982(11):85211 ENERGY
 TITLE: Airborne radiological sampling of Mount St. Helens plumes.
 AUTHOR: Andrews, V.E. [United States]
 CORPORATE SOURCE: Office of Radiation Programs, Las Vegas, NV (USA)
 (9511167)
 NUMBER OF REPORT: PB--81-213795
 DOCUMENT TYPE: Apr 1981. 20 p. Availability: NTIS, PC A02/MF A01.
 Report
 COUNTRY: United States
 LANGUAGE: English
 DOCUMENT NUMBER: ERA-07:034689

ABSTRACT: Particulate and gaseous samples for radiological analyses were collected from the plumes created by eruptions of Mount St. Helens. The sampling aircraft and equipment used are routinely employed in aerial radiological surveillance at the Nevada Test Site by the Environmental Protection Agency's Environmental Monitoring Systems Laboratory in Las Vegas, Nevada. An initial sample set was collected on April 4, 1980, during the period of recurring minor eruptions. Samples were collected again on May 19 and 20 following the major eruption of May 18. The Environmental Protection Agency's Office of Radiation Programs analyzed the samples for uranium and thorium isotopes, radium-226, lead-210, polonium-210, and radon-222. Other laboratories analyzed samples to determine particle size distribution and elemental composition. The only samples containing radioactivity above normal ambient levels were collected on May 20. Polonium-210 concentrations in the plume, determined from a sample collected between 5 and 30 km from the crater, were approximately an order of magnitude above background. Radon-222 concentrations in samples collected from the plume centerline at a distance of 15 km averaged approximately four times the average surface concentrations. The small increases in radioactivity would cause no observable adverse health effects.

CLASSIFICATION CODE: *500300; 510301
 CONTROLLED TERM: *LAVA; *CHEMICAL COMPOSITION; *VOLCANIC GASES:

*CHEMICAL COMPOSITION; *MT ST HELENS: *LAVA; *MT ST HELENS: *RADIATION MONITORING; *LAVA: *RADIATION MONITORING; *VOLCANIC GASES: *RADIATION MONITORING; *MT ST HELENS: *VOLCANIC GASES; AERIAL SURVEYING; AIR POLLUTION; DISTRIBUTION; LEAD; LEAD 210; PARTICLE SIZE; PLUMES; POLONIUM; POLONIUM 210; RADIOACTIVE AEROSOLS; RADIOACTIVE MATERIALS; RADIOACTIVITY; RADIUM; RADIUM 226; RADON; RADON 222; SAMPLING; THORIUM ISOTOPES; URANIUM ISOTOPES

BROADER TERM: ACTINIDE ISOTOPES; AEROSOLS; ALKALINE EARTH ISOTOPES; ALKALINE EARTH METALS; ALPHA DECAY RADIOISOTOPES; BETA DECAY RADIOISOTOPES; BETA-MINUS DECAY RADIOISOTOPES; COLLOIDS; DAYS LIVING RADIOISOTOPES; DISPERSIONS; ELEMENTS; EVEN-EVEN NUCLEI; FLUIDS; GASES; HEAVY NUCLEI; ISOTOPES; LEAD ISOTOPES; MATERIALS; METALS; MONITORING; MOUNTAINS; NONMETALS; NUCLEI; POLLUTION; POLONIUM ISOTOPES; RADIOISOTOPES; RADIUM ISOTOPES; RADON ISOTOPES; RARE GASES; SIZE; SOLS; YEARS LIVING RADIOISOTOPES

L84 ANSWER 28 OF 31 ENERGY COPYRIGHT 2008 USDOE/IEA-ETDE on STN

ACCESSION NUMBER: 1983(1):7074 ENERGY Full-text

TITLE: Focusing chromatography using a dilute precipitation reagent. Separation of 226Ra decayed nuclides by potassium fluoride.

AUTHOR: Furushima, K.; Shinagawa, M. (Kinki Univ., Higashi-Osaka, Osaka (Japan). Faculty of Science and Technology) [Japan]

SOURCE: Radioisotopes (Tokyo) (Jun 1981) v. 30(6) p. 299-304
CODEN: RAISAB ISSN: 0033-8303

DOCUMENT TYPE: Journal

COUNTRY: Japan

LANGUAGE: Japanese

ABSTRACT: The electrophoresis of 226Ra and it's decayed nuclides was carried out by using hydrochloric acid in the positive electrode cell and aqueous solution of potassium fluoride in the negative electrode cell. Experimental factors, i.e. concentrations of potassium fluoride and hydrochloric acid, intensity of electric field, duration for electrophoresis and pH value of the potassium fluoride solution etc. were examined. The sample solution was of 0.01 M hydrochloric acid with a tracer amount of 226Ra and 210Pb. Each of these nuclides was carrier free and was 4GBq/l(1muCi/ml) in concentration. For the sake of autoradiography, a photoengraving film (Fujilith Ortho Film, TAC = 135) was made use of examining the locations of the separated radioactive bands obtained on the paper strip. Their beta and gamma radioactivities were detected as the film darkening on developing the film. According to the positions thus detected, the strip filter paper was cut into pieces and the species of radionuclides were determined by the measurements of energies and half-lives on beta and gamma rays. The locations of the separated alpha-radionuclides were decided by etching the surface of it's film with 6 M aqueous solution of sodium hydroxide and the species of radionuclides were determined by the solid state alpha-track detection method. As the result, the optimum conditions for the separation are to use 0.01 M hydrochloric acid in the positive

CLASSIFICATION CODE: *400703; B1300

CONTROLLED TERM: *RADIA 226: *CHROMATOGRAPHY; AUTORADIOGRAPHY; DAUGHTER PRODUCTS; HYDROCHLORIC ACID; LEAD 210; PH VALUE; POTASSIUM FLUORIDES; PRECIPITATION; QUANTITY RATIO; ELECTROPHORESIS; AQUEOUS SOLUTIONS; ELECTRODES; BISMUTH 210; POLONIUM 210; LEAD 214

ELEMENT TERM: Ra; 226Ra; is; Ra is; Pb; 210Pb; Pb is

L84 ANSWER 29 OF 31 EMBASE COPYRIGHT (c) 2008 Elsevier B.V. All rights

reserved on STN

ACCESSION NUMBER: 1997217207 EMBASE Full-text

TITLE: Retrospective assessment of radon exposure by measurements of $(210)\text{Po}$ implanted in surfaces using an alpha track detector technique.

AUTHOR: Falk, R. (correspondence); Mellander, H.; Nyblom, L.; Ostergren, I.

CORPORATE SOURCE: Swed. Radiation Protection Institute, S-171 16 Stockholm, Sweden.

AUTHOR: Falk, R. (correspondence)

CORPORATE SOURCE: Swedish Radiation Protection Inst., S-171 16 Stockholm, Sweden.

SOURCE: Environment International, (1997) Vol. 22, No. SUPPL. 1, pp. S857-S861.

Refs: 4

ISSN: 0160-4120 CODEN: ENVIDV

PUBLISHER IDENT.: S 0160-4120(96)00193-6

COUNTRY: United Kingdom

DOCUMENT TYPE: Journal; Conference Article; (Conference paper)

FILE SEGMENT: 046 Environmental Health and Pollution Control

LANGUAGE: English

SUMMARY LANGUAGE: English

ENTRY DATE: Entered STN: 22 Aug 1997
Last Updated on STN: 22 Aug 1997

ABSTRACT: The radon exposure of the past is important in epidemiological studies where an assessment of lung cancer risk from indoor ***radon*** exposure is evaluated. The long-lived decay product, (210Po) Pb ($T_{1/2} = 22$ y), is implanted into indoor surfaces by alpha recoils and can be monitored to give information about the previous ***radon*** history. This gives an alternative or complementary method to the traditional measurements of the current average radon concentration. Autoradiographic alpha-track methods to assess the (210Po) Pb activity implanted in glass surfaces by measurement of (210Po) alpha activity were investigated to find a simple and reliable method for field use. One limiting factor at low exposure levels is the alpha background activity in the sub-surface material. In the search for a practical field method, the use of two different alpha-track detector materials was found successful. By exposing one Kodak LR-115 cellulose-nitrate ***film*** and one CR- 39 detector side by side on glass panes, the background of the glass is measured with the LR-115 and both the background and the signal by the CR-39 detector. Results from measurements in 31 dwellings show that an exposure of more than 1000 Bq.ovrhdot.y.ovrhdot.m(-3) to a glass surface can be measured with the (CR- LR) difference technique. Experiences from the field measurements show the method to be accurate, simple, and reliable and therefore a promising tool for future radon epidemiological studies.

CONTROLLED TERM: Medical Descriptors:
alpha radiation
autoradiography
*cancer risk
conference paper
*environmental exposure
priority journal
reliability
retrospective study

CONTROLLED TERM: Drug Descriptors:
*radon

CAS REGISTRY NO.: (radon) 10043-92-2

L84 ANSWER 30 OF 31 COMPENDEX COPYRIGHT 2008 EEI on STN
 ACCESSION NUMBER: 1984(11):201224 COMPENDEX Full-text
 DOCUMENT NUMBER: 8411121412
 ; *8491369
 TITLE: STUDY OF ^{210}Pb AND ^{210}Po
 DISTRIBUTIONS IN ENVIRONMENTAL SAMPLES BY CR-39 TRACK
 DETECTOR.
 AUTHOR: Hunyadi, I. (Hungarian Acad of Sciences, Inst of
 Nuclear Research, Debrecen, Hung); Somogyi, G.;
 Szilagyi, S.
 SOURCE: Nucl Tracks Radiat Meas v 8 n 1-4 1984, Solid State
 Nucl Track Detect, Proc of the Int Conf, 12th,
 Acapulco, Mex, Sep 4-10 1983 p 491-495
 SOURCE: Nucl Tracks Radiat Meas v 8 n 1-4 1984, Solid State
 Nucl Track Detect, Proc of the Int Conf, 12th,
 Acapulco, Mex, Sep 4-10 1983 p 491-495
 CODEN: NUTRDQ ISSN: 0191-278X

PUBLICATION YEAR: 1983
 LANGUAGE: English

ABSTRACT: Activity concentration distributions of long-lived alpha-emitters in aerosol samples are analyzed by high-resolution autoradiography in CR-39. A study of the alpha-activity attached to aerosols of different particulate sizes separated by a cascade impactor is also performed. It is found that, in the majority of samples, the alpha-activity can be dominantly related to the presence of Po-210 produced by its beta-active precursor Pb-210. Analysis of alpha-decay properties was done by autoradiographs taken at different post-sampling times. Spectroscopic studies of individual alpha tracks and track clusters were developed for high resolution alpha energy determination. Measured parameters were the major axis of surface track opening, the diameter of etched out track end, the total length measurable on the surface along the projected track, and the thickness of the layer etched away from the detector surface. 5 refs. CLASSIFICATION CODE: 944 Moisture, Pressure & Temperature, & Radiation

CONTROLLED TERM: Measuring Instruments; 443 Meteorology; 622
 Radioactive Materials; 815 Plastics & Polymeric
 Materials; 932 High Energy, Nuclear & Plasma Physics
 *PARTICLE DETECTORS: Applications; ATMOSPHERIC
 RADIOACTIVITY: Analysis; POLYCARBONATES
 SUPPLEMENTARY TERM: CR-39; AUTORADIOGRAPHY; ENVIRONMENTAL ALPHA ACTIVITY;
 SSNTD; SOLID STATE NUCLEAR TRACK DETECTORS
 ELEMENT TERM: Po; Pb; ^{210}Pb ; is; Pb is; ^{210}Po ; Po is

L84 ANSWER 31 OF 31 SCISEARCH COPYRIGHT(c)2008 The Thomson Corporation on STN
 ACCESSION NUMBER: 1994:283522 SCISEARCH Full-text

THE GENUINE ARTICLE: NJ812
 TITLE: AN EFFICIENT QUANTITATIVE TECHNIQUE FOR THE SIMULTANEOUS
 ANALYSES OF RADON DAUGHTERS PB-210 , BI-210 AND PO-210
 AUTHOR: CHURCH T M (Reprint); HUSSAIN N; FERDELMAN T G; FOWLER S W
 CORPORATE SOURCE: UNIV DELAWARE, COLL MARINE STUDIES, NEWARK, DE 19716
 (Reprint); IAEA, ENVIRONM LAB, MONACO 98012, MONACO
 COUNTRY OF AUTHOR: USA; MONACO
 SOURCE: TALANTA, (FEB 1994) Vol. 41, No. 2, pp. 243-249.
 ISSN: 0039-9140.
 PUBLISHER: ELSEVIER SCIENCE BV, PO BOX 211, 1000 AE AMSTERDAM,
 NETHERLANDS.
 DOCUMENT TYPE: Article; Journal
 FILE SEGMENT: PHYS
 LANGUAGE: English
 REFERENCE COUNT: 29
 ENTRY DATE: Entered STN: 1994

Last Updated on STN: 1994

ABSTRACT:

An improved and time efficient technique has been developed for quantitative determination of the long-lived Rn-222 daughters (Pb-210, ***Po*** -210 and Bi-209) in atmospheric and oceanic samples. The sample is first spiked with yield tracers for polonium (208 or 209), bismuth (207), and lead (stable lead carrier). These nuclides may then be scavenged through iron hydroxide precipitation and redissolved in a dilute (pH approximately 2) nitric acid plating medium with citrate and hydroxylamine hydrochloride at 90-degrees centigrade with constant stirring. First a silver planchet is suspended in the solution which plates polonium to high efficiency. Second, a nickel planchet is suspended in the same solution which is maintained hermetic (e.g. bubbling with helium) and bismuth is plated next with high efficiency. Third, lead is purified from the same solution using anion exchange techniques and isolated for beta counting as the sulfate. Polonium is analyzed by isotope dilution alpha spectrometry. Bismuth and lead are analyzed by anti-coincident beta counting in a low level shield. In the case of bismuth, the 207 tracer is added in quantities at least comparable to the background of the beta system such that counting before and after the decay of 2 Bi-210 gives the bismuth yield.

The unique characteristics of this technique are its speed and efficiency; all three radon daughters can be isolated for counting within 4 hr of pre-treating the sample. The remaining solution can be treated subsequently for other analyses as appropriate.

CATEGORY: CHEMISTRY, ANALYTICAL

SUPPL. TERM PLUS: GROUNDWATERS; SEPARATION; POLONIUM

REFERENCE(S):

| Referenced Author (RAU) | Year (R PY) | VOL (R VL) | ARN PG (R PG) | Referenced Work (RWK) |
|----------------------------|----------------|---------------|------------------|--------------------------|
| ANON | 1984 | 170 | 170 | EXPT NUCLEAR SCI |
| ANDREWS, J N | 1989 | 53 | 1791 | GEOCHIM COSMOCHIM AC |
| BAGNALL, K W | 1957 | 1 | 95 | CHEM RARE RADIOELEM |
| BLACK, S C | 1961 | 7 | 87 | HLTH PHYS |
| BLANCHARD, R L | 1966 | 38 | 189 | ANAL CHEM |
| CURIE, M | 1910 | 150 | 386 | CR HEBD ACAD SCI |
| GOLDBERG, E D | 1962 | 26 | 417 | GEOCHIM COSMOCHIM AC |
| GOLDBERG, E D | 1974 | 5 | 489 | SEA |
| HARADA, K | 1989 | 53 | 143 | GEOCHIM COSMOCHIM AC |
| HELKAMP, R W | 1979 | 30 | 237 | INT J APPL RADIAT IS |
| HUSSAIN, N | 1982 | 58 | 430 | EARTH PLANET SC LETT |
| HUSSAIN, N | 1980 | 44 | 1287 | GEOCHIM COSMOCHIM AC |
| HUSSAIN, N | 1984 | 1 | 208 | THESIS GUJARAT U |
| HUSSAIN, N | 1988 | 1 | 1 | 3RD CAMB SCH MIN Q R |
| JUNGE, C E | 1963 | 1 | 139 | AIR CHEM RADIOACTIVI |
| KHARKAR, D P | 1966 | 30 | 621 | GEOCHIM COSMOCHIM AC |
| KRISHNASWAMI, S | 1976 | 83 | 143 | ANAL CHIM ACTA |
| KRISHNASWAMI, S | 1982 | 18 | 1663 | WATER RESOUR RES |
| LAL, D | 1960 | 31 | 305 | REV SCI INSTRUM |
| LAMBERT, G | 1965 | 206 | 1343 | NATURE |
| MACKENZIE, A B | 1979 | 104 | 1151 | ANALYST |
| MARCKWALD, W | 1905 | 38 | 1591 | BER |
| MOORE, H E | 1973 | 78 | 17065 | J GEOPHYS RES |
| NARITA, H | 1989 | 36 | 925 | TALANTA |
| OSMOND, J K | 1982 | 1 | 1 | URANIUM SERIES DISEQ |
| POET, S E | 1972 | 77 | 16515 | J GEOPHYS RES |
| ROBBINS, J A | 1975 | 39 | 1285 | GEOCHIM COSMOCHIM AC |
| TUREKIAN, K K | 1977 | 5 | 1227 | ANNU REV EARTH PL SC |
| VONOETTINGEN, W F | 1930 | 10 | 1221 | PHYSIOL REV |

FILE 'HOME' ENTERED AT 14:09:55 ON 31 JUL 2008

SEARCH HISTORY

=> d his nofile

(FILE 'HOME' ENTERED AT 13:24:53 ON 31 JUL 2008)

FILE 'CAPLUS' ENTERED AT 13:25:06 ON 31 JUL 2008

E US2007-560922/APPS

E US2005-560922/APPS

L1 1 SEA ABB=ON US2005-560922/AP
D SCAN

FILE 'REGISTRY' ENTERED AT 13:26:01 ON 31 JUL 2008

L2 1 SEA ABB=ON 13981-52-7
L3 1 SEA ABB=ON 14255-04-0
D SCAN

FILE 'REGISTRY' ENTERED AT 13:26:55 ON 31 JUL 2008

D IDE L2

D IDE L3

FILE 'CAPLUS' ENTERED AT 13:27:14 ON 31 JUL 2008

L4 3024 SEA ABB=ON L2
L5 4376 SEA ABB=ON L3
L6 1219 SEA ABB=ON L4 AND L5
L7 28 SEA ABB=ON L2/P AND L3/P
E RADIATION SOURCES+ALL/CT
L8 2364 SEA ABB=ON RADIATION SOURCES/CT
L9 4 SEA ABB=ON L6 AND L8
D SCAN TI
L10 903425 SEA ABB=ON A/OBI
L11 118 SEA ABB=ON L10 AND L6
L12 6038 SEA ABB=ON L10(L) (SOURCE#/OBI OR EMIT#/OBI)
L13 28 SEA ABB=ON L12 AND L6
L14 1 SEA ABB=ON L12 AND L7
D SCAN
L15 258343 SEA ABB=ON SEAL#/BI
L16 2 SEA ABB=ON (L13 OR L7) AND L15
D SCAN TI
L17 881850 SEA ABB=ON FILM#/OBI
L18 76855 SEA ABB=ON POLYCARBONATE#/BI
L19 353488 SEA ABB=ON HYDROXIDE#/BI

FILE 'REGISTRY' ENTERED AT 13:32:37 ON 31 JUL 2008

E RADON/CN

L20 248 SEA ABB=ON RADON?/CN

FILE 'CAPLUS' ENTERED AT 13:32:52 ON 31 JUL 2008

L21 25986 SEA ABB=ON L20
L22 264 SEA ABB=ON L6 AND L21
L23 15 SEA ABB=ON (L7 OR L13) AND L21
D SCAN L1
L24 584884 SEA ABB=ON 71/SC,SX
L25 6 SEA ABB=ON L7 AND L21
L26 4 SEA ABB=ON L13 AND L21 AND L24
L27 0 SEA ABB=ON (L13 OR L7) AND L17
L28 3 SEA ABB=ON (L13 OR L7) AND L18
L29 2 SEA ABB=ON (L13 OR L7) AND L19
L30 3 SEA ABB=ON (L13 OR L7) AND (L17 OR L18 OR L19)
D AB L1

L31 1420531 SEA ABB=ON FILM#/BI
 L32 4 SEA ABB=ON (L13 OR L7) AND (L18 OR L19 OR L31)
 L33 5 SEA ABB=ON L4(L)PUR/RL
 L34 5 SEA ABB=ON L5(L)PUR/RL
 L35 2 SEA ABB=ON L33 AND L34
 D SCAN TI L33

INDEX '1MOBILITY, 2MOBILITY, ABI-INFORM, ADISCTI, AEROSPACE, AGRICOLA, ALUMINIUM, ANABSTR, ANTE, APOLLIT, AQUALINE, AQUASCI, AQUIRE, BABS, BIBLIODATA, BIOENG, BIOSIS, BIOTECHABS, BIOTECHDS, BIOTECHNO, CABA, CAOLD, CAPLUS, CASREACT, CBNB, CEABA-VTB, CERAB, ...' ENTERED AT 13:38:56 ON 31 JUL 2008

SEA (POLONIUM OR PO OR LEAD OR PB) (A) 210 OR PB210 OR 210PB OR 2

 29 FILE ABI-INFORM
 348 FILE AEROSPACE
 120 FILE AGRICOLA
 23 FILE ALUMINIUM
 320 FILE ANABSTR
 18 FILE ANTE
 530 FILE AQUALINE
 2100 FILE AQUASCI
 133 FILE AQUIRE
 100 FILE BABS
 27 FILE BIBLIODATA
 25 FILE BIOENG
 2400 FILE BIOSIS
 4 FILE BIOTECHABS
 4 FILE BIOTECHDS
 50 FILE BIOTECHNO
 692 FILE CABA
 214 FILE CAOLD
 5192 FILE CAPLUS
 1 FILE CASREACT
 20 FILE CEABA-VTB
 2 FILE CERAB
 11 FILE CIN
 288 FILE CIVILENG
 977 FILE COMPENDEX
 6 FILE COMPUAB
 159 FILE CONFSCI
 2 FILE COPPERLIT
 7 FILE CORROSION
 12 FILE CROPU
 14 FILE CSNB
 31 FILE DDFB
 9 FILE DDFU
 250 FILE DISSABS
 31 FILE DRUGB
 11 FILE DRUGU
 17 FILE ELCOM
 7 FILE EMA
 18 FILE EMBAL
 1570 FILE EMBASE
 64 FILE ENCOMPLIT
 4 FILE ENCOMPPAT
 5527 FILE ENERGY
 380 FILE ENVIROENG
 125 FILE EPFULL
 207 FILE ESBIOBASE

```

3   FILE FRANCEPAT
22  FILE FRFULL
12  FILE FROSTI
24  FILE FSTA
39  FILE GBFULL
52  FILE GENBANK
3560 FILE GEOREF
144 FILE HEALSAFE
95  FILE IFIPAT
5384 FILE INIS
84   FILE INPADOCDB
62   FILE INPAFAMDB
1831 FILE INSPEC
147 FILE INSPHYS
1   FILE IPA
3   FILE ITRD
13  FILE JAPIO
10  FILE KOREPAT
425 FILE LIFESCI
2   FILE MATBUS
148 FILE MECHENG
925 FILE MEDLINE
118 FILE METADEX
1   FILE NAPRALERT
43   FILE NLDB
941 FILE NTIS
756 FILE OCEAN
3   FILE PAPERCHEM2
1618 FILE PASCAL
4   FILE PATDPA
30  FILE PATDPAFULL
9   FILE PCI
236 FILE PCTFULL
1   FILE PHIN
4   FILE PIRA
1001 FILE POLLUAB
76  FILE PROMT
2   FILE RAPRA
3   FILE RSWB
2   FILE RUSSIAPAT
3578 FILE SCISEARCH
38  FILE SOLIDSTATE
1   FILE SOLIS
55  FILE TEMA
4517 FILE TOXCENTER
1   FILE TRIBO
102 FILE TULSA
22  FILE TULSA2
46  FILE UFORDAT
117 FILE ULIDAT
1110 FILE USPATFULL
333 FILE USPATOLD
145 FILE USPAT2
1271 FILE WATER
124 FILE WPIDS
124 FILE WPINDEX
7   FILE WSCA
QUE ABB=ON (POLONIUM OR PO OR LEAD OR PB) (A) 210 OR PB210 OR
210PB OR 210PO OR PO210
-----

```

D RANK

FILE 'STNGUIDE' ENTERED AT 13:40:57 ON 31 JUL 2008

FILE 'PASCAL, BIOSIS, GEOREF, ENERGY, DISSABS, CONFSCI, INSPEC, EMBASE, COMPENDEX, SCISEARCH' ENTERED AT 13:51:54 ON 31 JUL 2008

L37 7713 SEA ABB=ON (POLONIUM OR PO)(A) 210 OR 210PO OR PO210
 L38 17263 SEA ABB=ON (LEAD OR PB)(A) 210 OR 210PB OR PB210
 L39 2217612 SEA ABB=ON FILM#
 L40 68820 SEA ABB=ON RADON OR 222RADON OR RADON222
 L41 98370 SEA ABB=ON ALPHA(2A)(SOURCE OR EMIT? OR PARTICLE#)
 L42 47252 SEA ABB=ON POLYCARBONATE# OR POLY CARBONATE#
 L43 190082 SEA ABB=ON HYDROXIDE#
 L44 226644 SEA ABB=ON SEAL?
 L45 898 SEA ABB=ON L37 AND L38 AND (L39 OR L40 OR L41 OR L42 OR L43 OR L44)
 L46 14 SEA ABB=ON L37 AND L38 AND L39
 L47 694 SEA ABB=ON L37 AND L38 AND L40
 L48 252 SEA ABB=ON L37 AND L38 AND L41
 L49 2 SEA ABB=ON L37 AND L38 AND L42
 L50 24 SEA ABB=ON L37 AND L38 AND L43
 L51 21 SEA ABB=ON L37 AND L38 AND L44
 L52 57 SEA ABB=ON L37 AND L38 AND (L39 OR L42 OR L43 OR L44)
 L53 35 DUP REM L52 (22 DUPLICATES REMOVED)
 ANSWER '1' FROM FILE PASCAL
 ANSWERS '2-11' FROM FILE BIOSIS
 ANSWERS '12-29' FROM FILE ENERGY
 ANSWERS '30-31' FROM FILE EMBASE
 ANSWERS '32-34' FROM FILE COMPENDEX
 ANSWER '35' FROM FILE SCISEARCH
 L54 9 SEA ABB=ON L37 AND L38 AND L39 AND ((L40 OR L41 OR L43 OR L44))
 L55 94 SEA ABB=ON L37 AND L38 AND L40 AND ((L41 OR L43 OR L44))
 L56 6 SEA ABB=ON L37 AND L38 AND L43 AND ((L40 OR L41 OR L44))
 L57 4 SEA ABB=ON L37 AND L38 AND L44 AND ((L40 OR L41))
 L58 2389 SEA ABB=ON L37(2A) L38
 L59 501 SEA ABB=ON L58 AND (L40 OR L41)
 L60 36 SEA ABB=ON L58 AND L40 AND L41
 L61 907 SEA ABB=ON L40(5A) COLLECT?
 L62 0 SEA ABB=ON L58 AND L61 AND L41
 L63 1 SEA ABB=ON L58 AND L61
 D SCAN
 L64 29 DUP REM L60 (7 DUPLICATES REMOVED)
 ANSWER '1' FROM FILE PASCAL
 ANSWER '2' FROM FILE BIOSIS
 ANSWERS '3-18' FROM FILE ENERGY
 ANSWERS '19-26' FROM FILE INSPEC
 ANSWER '27' FROM FILE EMBASE
 ANSWERS '28-29' FROM FILE SCISEARCH
 L65 36 SEA ABB=ON L60 NOT (L49 OR L54 OR L56 OR L57 OR L63)
 D SCAN L64
 L66 1616 SEA ABB=ON RANDOM PULS?
 L67 3 SEA ABB=ON L60 AND L66
 D QUE
 L68 5 SEA ABB=ON L58 AND (L40 OR L41) AND L66

FILE 'WPIX' ENTERED AT 14:02:25 ON 31 JUL 2008

L69 1 SEA ABB=ON MITSUGASHIRA H?/AU
 D SCAN
 D TRIAL

| | | |
|-----|--------|--|
| | | E K08-A+ALL/MC |
| L70 | 68 | SEA ABB=ON (POLONIUM/BI, ABEX OR PO/BI, ABEX) (A) 210/BI, ABEX OR 210PO/BI, ABEX OR PO210/BI, ABEX OR POLONIUM210/BI, ABEX OR 210POLONIUM/BI, ABEX |
| L71 | 68 | SEA ABB=ON (LEAD/BI, ABEX OR PB/BI, ABEX) (A) 210/BI, ABEX OR 210PB/BI, ABEX OR PB210/BI, ABEX OR 210LEAD/BI, ABEX OR LEAD210/BI, ABEX |
| L72 | 11 | SEA ABB=ON L70 AND L71 |
| | | D SCAN |
| L73 | 974355 | SEA ABB=ON FILM#/BI, ABEX |
| L74 | 209521 | SEA ABB=ON ALPHA/BI, ABEX |
| L75 | 66573 | SEA ABB=ON POLYCARBONATE#/BI, ABEX OR POLY CARBONATE#/BI, ABEX |
| L76 | 146392 | SEA ABB=ON HYDROXIDE#/BI, ABEX |
| L77 | 722352 | SEA ABB=ON SEAL?/BI, ABEX |
| L78 | 1259 | SEA ABB=ON RANDOM PULS?/BI, ABEX |
| L79 | 9 | SEA ABB=ON L70 AND L71 AND (L73 OR L74 OR L75 OR L76 OR L77 OR L78) |
| L80 | 2652 | SEA ABB=ON ALPHA/BI, ABEX (2A) (SOURCE/BI, ABEX OR EMIT?/BI, ABEX OR PARTICLE#/BI, ABEX) |
| L81 | 4 | SEA ABB=ON L70 AND L71 AND (L73 OR L75 OR L76 OR L77 OR L78 OR L80) |

FILE 'STNGUIDE' ENTERED AT 14:07:22 ON 31 JUL 2008
FILE 'CAPLUS' ENTERED AT 14:09:11 ON 31 JUL 2008

D QUE L9
D QUE L16
D QUE L25
D QUE L26
D QUE L32
D QUE L35

L82 14 SEA ABB=ON (L9 OR L16 OR L25 OR L26 OR L32 OR L35)

FILE 'WPIX' ENTERED AT 14:09:14 ON 31 JUL 2008
D QUE L81

AL, BIOSIS
SCISEARCH
D QUE L49
D QUE L54
D QUE L56
D QUE L57
D QUE L63
D QUE L68

1.83 23 SEA ABB=ON (1.49 OB 1.54 OB 1.56 OB 1.57 OB 1.63 OB 1.68)

FILE 'CAPLUS, WPIX, BIOSIS, ENERGY, INSPEC, EMBASE, COMPENDEX, SCISEARCH'
ENTERED AT 14:09:22 ON 31 JUL 2008

ENTERED AT 11:05:22 ON 31-JUL-2005
31 DUP REM L:82 L:81 L:83 (10 DUPLICATES REMOVED)

ANSWERS '1-14' FROM FILE CAPLUS
ANSWERS '15-16' FROM FILE WPIX
ANSWERS '17-20' FROM FILE BIOSIS
ANSWERS '21-28' FROM FILE ENERGY
ANSWER '29' FROM FILE EMBASE
ANSWER '30' FROM FILE COMPENDEX
ANSWER '31' FROM FILE SCISEARCH
D IBIB AB HITIND 1-14
D IALL ABEX TECH 15-16
D IALL 17-31

FILE 'HOME' ENTERED AT 14:09:55 ON 31 JUL 2008